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<b>(21) International Application Number:</b> PCT/US99/17282 <b>(22) International Filing Date:</b> 29 July 1999 (29.07.99) <b>(30) Priority Data:</b> 60/094,690 30 July 1998 (30.07.98) US <b>(63) Related by Continuation (CON) or Continuation-in-Part (CIP) to Earlier Application</b> US 60/094,690 (CIP) Filed on 30 July 1998 (30.07.98) <b>(71) Applicant (for all designated States except US):</b> THE GOVERNMENT OF THE UNITED STATES OF AMERICA, as represented by THE SECRETARY OF HEALTH AND HUMAN SERVICES, NATIONAL INSTITUTES OF HEALTH [US/US]; Office of Technology Transfer, Suite 325, 6011 Executive Boulevard, Rockville, MD 20852-3804 (US). <b>(72) Inventors; and</b> <b>(75) Inventors/Applicants (for US only):</b> KLEINMAN, Hynda, K. [US/US]; National Institute of Dental and Craniofacial Research NIH, Building 30, Room 433, 30 Convent Drive, MSC-4370, Bethesda, MD 20892-4370 (US).		<p>GOLDSTEIN, Allan [US/US]; George Washington University, Dept. of Biochemistry, 2300 I Street, N.W., Washington, DC 20037 (US). MALINDA, Katherine, M. [US/US]; National Institute of Dental and Craniofacial Research NIH, Building 30, Room 433, 30 Convent Drive, MSC-4370, Bethesda, MD 20892-4370 (US). SOSNE, Gabriel [US/US]; 25341 Ronald Court, Oak Park, MI 48237 (US).</p> <p><b>(74) Agent:</b> WETHERELL, John, R. Jr.; Fish &amp; Richardson P.C., Suite 1400, 4225 Executive Square, La Jolla, CA 92037 (US).</p> <p><b>(81) Designated States:</b> AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).</p> <p><b>Published</b> <i>With international search report.</i> <i>Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i></p>
<b>(54) Title:</b> THYMOSIN $\beta$ 4 PROMOTES WOUND REPAIR <b>(57) Abstract</b> <p>The present invention relates to methods for promoting tissue repair, angiogenesis and cell migration. The method of the invention utilizes thymosin <math>\beta</math>4 (T<math>\beta</math>4) peptide to promote tissue repair, angiogenesis and cell migration. The invention further relates to modulating T<math>\beta</math>4 activity in tissues.</p>		

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**THYMOSIN  $\beta$ 4 PROMOTES WOUND REPAIR****STATEMENT AS TO FEDERALLY SPONSORED RESEARCH**

This invention was made in part with funds from the National Institutes of  
5 Health, Intramural Program. The government may have certain rights in this invention.

**CROSS REFERENCE TO RELATED APPLICATIONS**

This application claims priority from Provisional Application Serial No.  
60/094,690, filed July 30, 1998, which is incorporated herein by reference in its  
entirety and to which application a priority claim is made under 35 U.S.C. §119(e).

10

**TECHNICAL FIELD OF THE INVENTION**

The present invention relates generally to tissue repair and more  
specifically to methods of wound healing using thymosin  $\beta$ 4.

**BACKGROUND OF THE INVENTION**

Inadequate methods and compositions to effectively heal chronic wounds  
15 is a significant health care problem. Impaired wound healing increases the chances of  
mortality and morbidity. This problem is especially prominent in patients with  
diabetes who develop severe, life threatening wounds on body extremities. Chronic  
diabetic foot ulcers often lead to amputations. These wounds are often the result of  
poor circulation derived from the diabetic patients' insulin-compromised cells as well  
20 as impaired vascularization of the wound bed, reduced infiltration of germ fighting  
cells and reduced tissue epithelialization. As a result, most current therapies include  
attempts to revascularize the wound bed and prevent infection.

Wounds in non-compromised tissues undergo a complex and ordered  
series of events to repair the tissue. The series of events may include infiltration of  
25 immune cells as part of the process to remove and destroy necrotic tissue, increased  
vascularization by angiogenic factors and increased cell proliferation and extracellular  
matrix deposition. Although the basic process of tissue repair has been characterized,

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the individual steps and factors necessary to carry out this complex series of events are not well understood. The identification of individual steps and factors could lead to improved methods for the treatment of diseases resulting from inadequate wound repair processes.

5           Previous studies have used the "scratch" wound closure assay to assess the potential effects of an agent on *in vitro* cell migration. Though informative, such a test does not mimic the dynamic *in vivo* wound healing conditions to the extent that not all factors involved in wound closure are present in the *in vitro* assay. For this reason, *in vivo* systems have been developed to assess the ability of an agent or factor  
10 to modulate wound healing activities.

          Using these types of *in vitro* models, a number of specific growth factors have been recognized for their effect on angiogenesis. One such growth factor is TGF- $\beta$ . This family of dimeric proteins includes TGF- $\beta$ 1, TGF- $\beta$ 2, TGF- $\beta$ 3, TGF- $\beta$ 4, and TGF- $\beta$ 5 which regulate the growth and differentiation of many cell types.  
15 This family of proteins exhibits a range of biological effects from stimulating the growth of some cell types (Noda *et al.*, (1989) *Endocrinology*, 124:2991-2995) and inhibiting the growth of other cell types (Goey *et al.*, (1989) *J. Immunol.*, 143:877-880; Pietenpol *et al.*, (1990) *Proc. Nat'l. Acad. Sci. USA*, 87:3758-3762). TGF- $\beta$  has also been shown to increase the expression of extracellular matrix proteins, including  
20 collagen and fibronectin (Ignatz *et al.*, (1986) *J. Biol. Chem.*, 261:4337-4345) and accelerates the healing of wounds (Mustoe *et al.*, (1987) *Science*, 237:1333-1335).

          Another growth factor recognized for its effect on angiogenesis is Platelet Derived Growth Factor (PDGF). PDGF was originally found to be a potent mitogen for mesenchymal derived cells (Ross R. *et al.* (1974) *Proc Nat'l Acad Sci USA*  
25 71(4):1207-1210.; Kohler N. *et al.* (1974) *Exp. Cell Res.* 87:297-301). Further studies have shown that PDGF increases the rate of cellularity and granulation in tissue formation. Wounds treated with PDGF have the appearance of an early stage inflammatory response, including an increase in neutrophils and macrophage cell types at the wound site. These wounds also show enhanced fibroblast function  
30 (Pierce, GF *et al.* (1988) *J. Exp. Med.* 167:974-987). Both PDGF and TGF $\beta$  have

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been shown to increase collagen formation, DNA content, and protein levels in animal studies. (Grotendorst, GR *et al.* (1985) *J. Clin. Invest.* 76:2323-2329.; Sporn, MB *et al.* (1983) *Science* 219:1329). The effect of PDGF in wound healing has been shown to be effective in human wounds. In human wounds, PDGF-AA expression is

5 increased within pressure ulcers undergoing healing. The increase of PDGF-AA corresponds to an increase in activated fibroblasts, extracellular matrix deposition, and active vascularization of the wound. Furthermore, such an increase in PDGF-AA is not seen in chronic non-healing wounds. A number of other growth factors having the ability to induce angiogenesis and wound healing include, Vascular Endothelial

10 Growth Factor (VEGF), Keratinocyte Growth Factor (KGF) and basic Fibroblast Growth Factor (bFGF).

However, most of these growth and angiogenic factors have side effects. Accordingly, there is a need for additional factors useful in promoting wound repair.

### SUMMARY OF THE INVENTION

15 The present invention is based on the discovery that thymosin  $\beta$ 4 (T $\beta$ 4) accelerates wound healing and stimulates wound repair. Based on this finding, it is now possible to develop methods for accelerating wound healing in subjects having wounds in need of such treatment.

In a first embodiment, the invention provides a method for promoting

20 wound repair in a subject in need of such treatment by administering to the subject or contacting the site of the wound with a wound-healing effective amount of a composition containing a wound healing polypeptide comprising the amino acid sequence LKKTET and conservative variants thereof having wound healing activity. In one aspect of the method, the wound healing polypeptide is T $\beta$ 4 or an isoform of

25 T $\beta$ 4.

In another embodiment, the invention provides a method for promoting tissue repair in a tissue in need of such treatment by contacting the tissue with an effective amount of a composition containing a wound healing polypeptide comprising the amino acid sequence LKKTET and conservative variants thereof

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having wound healing activity, or nucleic acid encoding a wound healing polypeptide.

In one aspect of the method, a wound healing peptide is T $\beta$ 4 or an isoform of T $\beta$ 4.

The tissue may be contacted either *in vivo* or *ex vivo*.

In yet another embodiment, the invention provides a method of modulating  
5 wound repair in a subject in need of such treatment by systemic delivery of a wound-healing effective amount of a wound healing polypeptide comprising the amino acid sequence LKKTET and conservative variants thereof having wound healing activity. In one aspect of the method, a wound healing peptide is T $\beta$ 4 or an isoform of T $\beta$ 4.

In yet another embodiment, the present invention provides a method for  
10 stimulating epithelial cell migration at the site of a wound by contacting the wound with an effective amount of a T $\beta$ 4 polypeptide.

In another embodiment, the invention provides a method of diagnosing a pathological condition in a subject characterized by a wound healing disorder associated with T $\beta$ 4, including obtaining a sample suspected of containing T $\beta$ 4 from  
15 the subject, detecting a level of T $\beta$ 4 in the sample and comparing the level of T $\beta$ 4 with the level found in a normal sample (*i.e.*, a standard sample).

In another embodiment, the invention provides a method of ameliorating a wound healing disorder associated with T $\beta$ 4, including treating a subject having the disorder with a composition which modulates T $\beta$ 4 activity or the activity of a T $\beta$ 4  
20 isoform.

In yet another embodiment, the present invention provides pharmaceutical compositions comprising a wound healing polypeptide comprising the amino acid sequence LKKTET and conservative variants thereof having wound healing activity and a pharmaceutically acceptable carrier. In one aspect, the wound healing  
25 polypeptide is T $\beta$ 4 or an isoform of T $\beta$ 4.

The details of one or more embodiments of the invention are set forth in the accompanying drawings and the description below. Other features, objects, and advantages of the invention will be apparent from the description and drawings, and from the claims.

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### DESCRIPTION OF DRAWINGS

FIG. 1 is a schematic drawing of a wound.

FIG. 2 is a bar graph which shows the effect of topical and systemic delivery of T $\beta$ 4 on the width of a punch wound as compared to control. (A) Topical delivery of 5  $\mu$ g/50  $\mu$ l was performed on three of the six wounds in each animal on the day of wounding and at 48 hours after wounding. (B) Intraperitoneal injections of 60  $\mu$ g/300 $\mu$ l were done on the day of the wounding and thereafter every other day. Control animals were treated similarly with saline. Measurements are expressed as the mean percent decrease  $\pm$  SEM.

FIG. 3 is a bar graph which shows the effect of topical and systemic delivery of T $\beta$ 4 on the gap of a punch wound as compared to control. (A) Topical delivery of 5  $\mu$ g/50  $\mu$ l was performed on the day of wounding and at 48 hours after wounding. (B) Intraperitoneal injections of 60  $\mu$ g/300 $\mu$ l were done on the day of the wounding and thereafter every other day. Measurements are expressed as the mean percent decrease  $\pm$  SEM.

FIG. 4 is a histological section, stained with H&E, demonstrating the appearance of control and thymosin  $\beta$ 4 treated wounds at low magnification and higher magnification. Wounds are from day 7 as described in the legend to figure 2. Arrows indicate the edges of the original wound. (A) Control wound treated with saline. Migration of the epithelium is visible at the wound edges and debris are visible over the unhealed wound. (B) Increased re-epithelialization of the wound occurred when T $\beta$ 4 was injected intraperitoneally (60  $\mu$ g/300 $\mu$ l on alternate days). (C) Topical treatment (5 $\mu$ g/50 $\mu$ l of T $\beta$ 4) resulted in complete reepithelialization of the wound epidermis. Boxed areas are the location of the higher magnification fields (D-F). (D-F) Dermis near dermal and epidermal junction. (D) Control showing few cells near the dermis and little neovascularization. (E) and (F) Dermis showing granulation tissue infiltrated with fibroblasts and extensive neovascularization (arrowheads). (E) Intraperitoneal treatment and (F) topical application both resulted in significant new capillaries. (Scale bar = 1 mm).

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FIG. 5 shows histological sections of 7 day wounds showing collagen deposition/accumulation. Masson's trichrome staining shows collagen and endothelial cells. (A) Low magnification view of a control wound treated with saline. (B) and (C). Low magnification views of wounds where T $\beta$ 4 was injected intraperitoneally (B) or applied topically (A). Boxed areas are the location of the higher magnification fields (D-F). Arrows indicate the edges of the original wound. (D) Control wound at higher magnification showing baseline collagen accumulation. Treatment intraperitoneally (E) or (F) topically resulted in enhanced collagen production/accumulation compared to wounds treated with saline. (Scale bar = 1 mm).

FIG. 6 shows T $\beta$ 4 stimulated keratinocyte migration in Boyden chamber assays. (A) T $\beta$ 4 in the lower wells of the chamber resulted in a 2-3 fold increase in migration on filters coated with collagen IV. The positive control, conditioned media, also showed increased migration over media alone.

FIG. 7 shows a graph demonstrating the migration of corneal epithelial cells at various concentrations of T $\beta$ 4.

FIG. 8 shows a graph representing corneal re-epithelialization in rat corneas in the presence and absence of T $\beta$ 4.

FIG. 9 shows a graph representing corneal re-epithelialization in the presence and absence of various concentrations of T $\beta$ 4.

FIG. 10 shows an amino acid sequence of T $\beta$ 4.

FIG. 11 shows the amino acid sequence of several known isoforms of T $\beta$ 4, and their phylogenetic distribution. N-terminal acetylation is indicated by "ac." Residues between 13 and 24 are thought to be important for actin binding.

## DETAILED DESCRIPTION OF THE INVENTION

Thymosin  $\beta$ 4 was initially identified as a protein that is up regulated during endothelial cell migration and differentiation *in vitro*. Thymosin  $\beta$ 4 was originally isolated from the thymus and is a 43 amino acid, 4.9 kDa ubiquitous polypeptide identified in a variety of tissues. Several roles have been ascribed to this



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protein including a role in endothelial cell differentiation and migration, T cell differentiation, actin sequestration and vascularization. One biological activity of thymosin  $\beta$ 4 (T $\beta$ 4), as shown herein, effects tissue repair and wound healing. Another activity of T $\beta$ 4 is anti-inflammatory activity.

5           The present invention resulted from investigation of the effects of T $\beta$ 4 on wound healing. *In vivo* results have demonstrated that topical and systemic delivery of T $\beta$ 4 promotes wound healing. Additional experiments demonstrated that T $\beta$ 4-treated wounds have increased extracellular matrix deposition in the wound bed.

          The present invention identifies T $\beta$ 4 as an active factor in promoting  
10   wound closure and tissue repair *in vivo* as well as increasing epithelial cell migration. *In vivo* administration of T $\beta$ 4 indicates that cell migration, angiogenesis and extracellular matrix deposition are stimulated at or above the levels observed for migration, angiogenesis and matrix deposition in control animals. T $\beta$ 4 promotes wound closure when administered systemically (*e.g.*, intra-peritoneally) and topically  
15   in wounded animal models. Increased levels of collagen were also observed in treated wounds showing that T $\beta$ 4 treatment can also accelerate wound contraction and stimulate the healing process.

          The methods of the invention result from the identification of the effect of T $\beta$ 4 on wound healing. *In vivo*, T $\beta$ 4 stimulates wound healing in a full thickness  
20   punch wound (see Example 1) and in repair of eye-related wounds (Example 4). When given either topically or systemically (*e.g.*, intra-peritoneally) T $\beta$ 4 accelerated closure and healing of wounds (see Example 1, 4, and 5).

#### **Promoting Tissue Regeneration**

          In one embodiment, the invention provides a method for accelerating  
25   wound healing in a subject by contacting a wound with a wound-healing effective amount of a composition which contains T $\beta$ 4 or a T $\beta$ 4 isoform. The contacting may be topically or systemically. Examples of topical administration include, for example, contacting the wound with a lotion, salve, gel, cream, paste, spray, suspension, dispersion, hydrogel, ointment, or oil comprising T $\beta$ 4. Systemic administration

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includes, for example, intravenous, intraperitoneal, intramuscular injections of a composition containing T $\beta$ 4 or a T $\beta$ 4 isoform. A subject may be any mammal, preferably human.

In addition, T $\beta$ 4 or a T $\beta$ 4 isoform is therapeutically valuable in cases  
5 where there is an impaired wound healing process, such as in wound healing compromised subjects. By "wound healing compromised" is meant subjects which have a reduced, decreased, or inability to recover from a wounding or trauma, due to recurrent wounding, trauma or inability of the subject's natural system to properly effectuate wound healing. For example, steroids reduce the ability of a subject to heal  
10 as compared to a subject which is not on steroids. Other such wounds present in compromised subjects include, but are not limited to, skin wounds such as diabetic ulcers, venus ulcers or pressure ulcers. Additionally, T $\beta$ 4 or a T $\beta$ 4 isoform is therapeutically valuable to augment the normal healing process.

As used herein, a "wound-healing effective amount" of a composition  
15 containing T $\beta$ 4 or a T $\beta$ 4 isoform for use in wound healing is defined as that amount that is effective in promoting tissue regeneration and repair. The "wound-healing effective amount" may be the therapeutically effective amount. Diseases, disorders or ailments possibly modulated by T $\beta$ 4 or a T $\beta$ 4 isoform include tissue repair subsequent to traumatic injuries or conditions including arthritis, osteoporosis and  
20 other musculo-skeletal disorders, burns, ulcers and other skin lesions, neurological and nerve disease and cardiovascular diseases including ischemia and atherosclerosis. Other potential tissues which can be treated by the methods and compositions of the invention include epidermal, eye, uro-genital, gastro-intestinal, cardiovascular, muscle, connective, and neural tissues. The term "induce", "induction" or "effect" as  
25 used herein, refers to the activation, stimulation, enhancement, initiation and/or maintenance of cellular mechanisms or processes necessary for the formation of a tissue or a portion thereof, repair process or tissue development as described herein.

### Modulation of Wound Healing

Wound healing, tissue regeneration and tissue repair result from a complex process that includes the proliferation and migration of inflammatory cells, endothelial cells, stromal cells and parenchymal cell, the deposition of extracellular matrix materials and the growth of new blood vessels, particularly capillaries. This  
5 complex process plays a crucial role in such beneficial functions as embryogenesis, the female reproductive cycle, as well as such abnormal functions as arthritis, chronic ulcerations and neuro-degenerative diseases.

In another embodiment, the invention provides a method for modulating  
10 wound healing in a subject or a tissue including contacting the subject or tissue with an effective wound-healing amount of a composition containing T $\beta$ 4 or a T $\beta$ 4 isoform. It is envisioned that T $\beta$ 4 or a T $\beta$ 4 isoform can be administered topically or systemically to prevent or treat a damaged tissue including, for example, tissues  
15 damaged due to ischemia, including ischemic brain, bone and heart disease, damage to corneal or retinal tissue of the eye, and damage to epithelial tissue, including skin.

In addition, the method of the invention is useful in promoting wound healing in tissues by promoting angiogenesis in tissue deprived of adequate blood flow. For example, a composition containing T $\beta$ 4 can promote the healing of chronic ulcers by increasing blood supply to the tissue site as well as increasing keratinocyte  
20 migration to close a wound.

T $\beta$ 4 isoforms have been identified and have about 70%, or about 75%, or about 80% or more homology to the amino acid sequence of T $\beta$ 4 set forth in Fig. 10. Such isoforms include, for example, T $\beta$ 4<sup>ala</sup>, T $\beta$ 9, T $\beta$ 10, T $\beta$ 11, T $\beta$ 12, T $\beta$ 13, T $\beta$ 14 and T $\beta$ 15 (Fig. 11; see also, Mihelic *et al.*, (1994) *Amino Acids*, 6:1-13, which describes  
25 the amino acid sequence of other T $\beta$ 4 isoforms, and is incorporated herein by reference). Similar to T $\beta$ 4, the T $\beta$ 10 and T $\beta$ 15 isoforms have been shown to sequester actin. T $\beta$ 4, T $\beta$ 10 and T $\beta$ 15, as well as these other isoforms share an amino acid sequence, LKKTET, that appears to be involved in mediating actin sequestration or binding. Although not wishing to be bound to any particular theory, the wound  
30 healing activity of T $\beta$ 4 and T $\beta$ 4 isoforms may be due, in part, to the ability to

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polymerize actin. For example, T $\beta$ 4 can modulate actin polymerization in wounds to promote healing (*e.g.*,  $\beta$ -thymosins appear to depolymerize F-actin by sequestering free G-actin). T $\beta$ 4's ability to modulate actin polymerization may therefore be due to all, or in part, its ability to bind to or sequester actin via the LKKTET sequence.

- 5 Thus, as with T $\beta$ 4, other proteins which bind or sequester actin, or modulate actin polymerization, including T $\beta$ 4 isoforms having the amino acid sequence LKKTET, are likely to promote wound healing alone, or in a combination with T $\beta$ 4, as set forth herein.

- Thus, it is specifically contemplated that known T $\beta$ 4 isoforms, such as
- 10 T $\beta$ 4<sup>ala</sup>, T $\beta$ 9, T $\beta$ 10, T $\beta$ 11, T $\beta$ 12, T $\beta$ 13, T $\beta$ 14, and T $\beta$ 15, as well as T $\beta$ 4 isoforms not yet identified, will be useful in the methods of the invention. As such T $\beta$ 4 isoforms are useful in the methods of the invention, including the methods practiced in a subject, the invention therefore further provides pharmaceutical compositions comprising T $\beta$ 4 isoforms T $\beta$ 4<sup>ala</sup>, T $\beta$ 9, T $\beta$ 10, T $\beta$ 11, T $\beta$ 12, T $\beta$ 13, T $\beta$ 14, and T $\beta$ 15 and
- 15 a pharmaceutically acceptable carrier.

- In addition, other proteins having actin sequestering or binding capability, or that can mobilize actin or modulate actin polymerization, as demonstrated in an appropriate sequestering, binding, mobilization or polymerization assay, or identified by the presence of an amino acid sequence that mediates actin binding, such as
- 20 LKKTET, for example, can similarly be employed in the methods of the invention. Such proteins include gelsolin, vitamin D binding protein (DBP), profilin, cofilin, depactin, DNaseI, vilin, fragmin, severin, capping protein,  $\beta$ -actinin and acumentin, for example. As such methods include those practiced in a subject, the invention further provides pharmaceutical compositions comprising gelsolin, vitamin D binding
- 25 protein (DBP), profilin, cofilin, depactin, DNaseI, vilin, fragmin, severin, capping protein,  $\beta$ -actinin and acumentin as set forth herein. Thus, the invention includes the use of wound healing polypeptide comprising the amino acid sequence LKKTET and conservative variants thereof.

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As used herein, the term "conservative variant" or grammatical variations thereof denotes the replacement of an amino acid residue by another, biologically similar residue. Examples of conservative variations include the replacement of a hydrophobic residue such as isoleucine, valine, leucine or methionine for another, the replacement of a polar residue for another, such as the substitution of arginine for lysine, glutamic for aspartic acids, or glutamine for asparagine, and the like.

T $\beta$ 4 has been localized to a number of tissue and cell types and thus, agents which stimulate the production of T $\beta$ 4 can be added to a composition to effect T $\beta$ 4 production from a tissue and/or a cell. Agents that effect wound repair can also be included in such a composition to augment the wound healing process. Such agents include members of the family of growth factors, such as insulin-like growth factor (IGF-1), platelet derived growth factor (PDGF), epidermal growth factor (EGF), transforming growth factor beta (TGF- $\beta$ ), basic fibroblast growth factor (bFGF), thymosin  $\alpha$ 1 (T $\alpha$ 1) and vascular endothelial growth factor (VEGF). More preferably, the agent is transforming growth factor beta (TGF- $\beta$ ) or other members of the TGF- $\beta$  superfamily. T $\beta$ 4 compositions of the invention aid in wound healing by effectuating growth of the connective tissue through extracellular matrix deposition, cellular migration and vascularization of the wound bed.

Additionally, agents that assist or stimulate the wound healing process may be added to a composition along with T $\beta$ 4 or a T $\beta$ 4 isoform to further modulate the wound healing process. Such agents include angiogenic agents, growth factors, agents that direct differentiation of cells, agents that promote migration of cells and agents that stimulate the provision of extracellular matrix materials in the wound bed. For example, and not by way of limitation, T $\beta$ 4 or a T $\beta$ 4 isoform alone or in combination can be added in combination with any one or more of the following agents: VEGF, KGF, FGF, PDGF, TGF $\beta$ , IGF-1, IGF-2, IL-1, prothymosin  $\alpha$  and thymosin  $\alpha$ 1 in a wound-healing effective amount.

In another aspect, the invention is useful for repair of tissue resulting from injuries due to surgical procedures, irradiation, laceration, toxic chemicals, viral infections, bacterial infections or burns. Additionally, the invention is useful for

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revitalizing scar tissue resulting from any number of procedures, accidents or trauma. The term "scar tissue" means fibrotic or collagenous tissue formed during the healing of a wound or other morbid process. For example, T $\beta$ 4 can be included in a controlled release matrix which can be positioned in proximity to damaged tissue

5 thereby promoting regeneration, repair and/or revascularization of such tissue. The term "controlled release matrix" means any composition that allows for the release of a bioactive substance which is mixed or admixed therein. The matrix can be a solid composition, a porous material (such as a scaffold, mesh, or sponge), or a semi-solid, gel or liquid suspension containing bioactive substances. The term "bioactive

10 material" means any composition that modulates tissue repair when used in accordance with the method of the present invention. The bioactive materials or matrix can be introduced by means of injection, surgery, catheters or any other means suitable for modulating tissue repair.

It is envisioned that the methods and compositions of the invention can be

15 used to aid wound healing and repair in guided tissue regeneration (GTR) procedures. Such procedures are currently used by those skilled in the medical arts to accelerate wound healing. Typically, nonresorbable or bioabsorbable materials are used to accelerate wound healing by promoting the repopulation of the wound area with cells which form the architectural and structural matrix of the tissue. For example, the

20 methods and compositions of the invention can be used in aiding tissue repair or regeneration at an ulcer site in a human or other subject by placing a composition containing a bioresorbable polymer and T $\beta$ 4 at the site in need of tissue repair or regeneration such that the composition is effective for aiding tissue regeneration by releasing a wound-healing effective amount of T $\beta$ 4 at the site.

25 In another aspect, the invention is useful for the purposes of promoting tissue growth during the process of tissue engineering. As used herein, "tissue engineering" is defined as the creation, design, and fabrication of biological prosthetic devices, in combination with synthetic or natural materials, for the creation, augmentation or replacement of body tissues and organs. Thus, the present method

30 can be used to augment the design and growth of human tissues outside the body, for

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later implantation inside the body, or augment the design and growth of a tissue inside the body to repair or replace diseased or damaged tissue. For example, T $\beta$ 4 may be useful in promoting the growth of skin graft replacements which are used as a therapy in the treatment of burns and ulcers.

5           In another aspect of tissue engineering, T $\beta$ 4 can be included in external or internal devices containing human tissue designed to replace the function of a diseased internal tissue. This approach involves isolating cells from the body, placing them on or within a three-dimensional matrices and implanting the new system inside the body or using the system outside the body. The methods and compositions of the  
10           invention can be used and included in such matrices to promote the growth of tissues contained in the matrices. For example, T $\beta$ 4 can be included in a tissue engineered construct to promote the growth of the cells contained in the construct. It is envisioned that the method of the invention can be used to augment tissue repair, regeneration and engineering in endothelial cell-related products which may contain  
15           cartilage, cartilage-bone composites, bone, central nervous system tissues, muscle, liver, pancreatic islet (insulin-producing) cells, urogenital tissues, breast and tissues for gene therapy applications.

          The present invention further provides methods and compositions for modulating female reproductive tract function. Growth factors have been shown to  
20           play a role in cyclic mitosis and differentiation of endometrial cellular components, recruitment of macrophages in decidualizing the endometrium, endometrial-trophoblast interactions, early pregnancy maintenance, and endometrial functional regeneration. The term "modulate" as used herein, denotes a modification of an existing condition or biologic state. Modulation of a condition as defined herein,  
25           encompasses both an increase or a decrease in the determinants affecting the existing condition. For example, administration of T $\beta$ 4 could be used to augment uterine functions in a condition where the promotion of endothelial cell growth is desired. For example, the uterus may be treated with T $\beta$ 4 to promote the growth and development of placental membranes or endometrial growth or the repair of these  
30           tissue following tissue injury. Furthermore, treatment with T $\beta$ 4 may be used to

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promote and maintain a pregnancy by facilitating endometrial-trophoblast interaction. Alternatively, antagonist to T $\beta$ 4 could be administered to modulate conditions of excessive endometrial growth in which the level of T $\beta$ 4 is excessive in comparison to a normal biological condition. In addition, T $\beta$ 4 in combination with other agents,  
5 such as thymosin  $\alpha$ 1, may be desirable for treating disorders of the reproductive tract.

The therapeutic approaches described herein involve various routes of administration or delivery of reagents or compositions comprising the T $\beta$ 4 of the invention including any conventional administration techniques (for example, but not limited to, topical administration, local injection, inhalation, or systemic  
10 administration), to a subject with a wound or tissue in need of healing or repair. Administration of T $\beta$ 4, as described above, can accelerate wound healing, increase cell migration into a wound site, induce the formation of tissue repair or regeneration, or promote the growth and development of the endometrium. The reagent, formulation or composition may also be targeted to specific cells or receptors by any  
15 method described herein or by any method known in the art of delivering, targeting T $\beta$ 4 polypeptides and expressing genes encoding T $\beta$ 4. For example, the methods and compositions using or containing T $\beta$ 4 of the invention may be formulated into pharmaceutical compositions by admixture with pharmaceutically acceptable non-toxic excipients or carriers. Such compositions may be prepared for parenteral  
20 administration, particularly in the form of liquid solutions or suspensions in aqueous physiological buffer solutions; for oral administration, particularly in the form of tablets or capsules; or for intranasal administration, particularly in the form of powders, nasal drops, or aerosols. Sustained release compositions are also encompassed by the present invention. Compositions for other routes of  
25 administration may be prepared as desired using standard methods.

A composition of the invention containing T $\beta$ 4 may be conveniently administered in unit dosage form, and may be prepared by any of the methods well known in the pharmaceutical art, for example, as described in Remington's Pharmaceutical Sciences (Mack Pub. Co., Easton, PA, 1990). Formulations for  
30 parenteral administration may contain as common excipients sterile water or saline,



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polyalkylene glycols such as polyethylene glycol, oils of vegetable origin, hydrogenated naphthalenes, and the like. In particular, biocompatible, biodegradable lactide polymer, lactide/glycolide copolymer, or polyoxethylene-polyoxypropylene copolymers are examples of excipients for controlling the release of a compound of the invention *in vivo*. Other suitable parenteral delivery systems include ethylene-vinyl acetate copolymer particles, osmotic pumps, implantable infusion systems, and liposomes. Formulations for inhalation administration may contain excipients such as lactose, if desired. Inhalation formulations may be aqueous solutions containing, for example, polyoxyethylene-9-lauryl ether, glycocholate and deoxycholate, or they may be oily solutions for administration in the form of nasal drops. If desired, the compounds can be formulated as a gel to be applied intranasally. Formulations for parenteral administration may also include glycocholate for buccal administration.

The composition of the liposome is usually a combination of phospholipids, particularly high-phase-transition-temperature phospholipids, usually in combination with steroids, especially cholesterol. Other phospholipids or other lipids may also be used. The physical characteristics of liposomes depend on pH, ionic strength, and the presence of divalent cations.

Examples of lipids useful in liposome production include phosphatidyl compounds, such as phosphatidylglycerol, phosphatidylcholine, phosphatidylserine, phosphatidylethanolamine, sphingolipids, cerebroside, and gangliosides. Particularly useful are diacylphosphatidylglycerols, where the lipid moiety contains from 14-18 carbon atoms, particularly from 16-18 carbon atoms, and is saturated. Illustrative phospholipids include egg phosphatidylcholine, dipalmitoylphosphatidylcholine and distearoylphosphatidylcholine.

The targeting of liposomes has been classified based on anatomical and mechanistic factors. Anatomical classification is based on the level of selectivity, for example, organ-specific, cell-specific, and organelle-specific. Mechanistic targeting can be distinguished based upon whether it is passive or active. Passive targeting utilizes the natural tendency of liposomes to distribute to cells of the reticulo-endothelial system (RES) in organs which contain sinusoidal capillaries. Active

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targeting, on the other hand, involves alteration of the liposome by coupling the liposome to a specific ligand such as a monoclonal antibody, sugar, glycolipid, or protein, or by changing the composition or size of the liposome in order to achieve targeting to organs and cell types other than the naturally occurring sites of

5 localization.

The surface of the targeted delivery system may be modified in a variety of ways. In the case of a liposomal targeted delivery system, lipid groups can be incorporated into the lipid bilayer of the liposome in order to maintain the targeting ligand in stable association with the liposomal bilayer. Various linking groups can be  
10 used for joining the lipid chains to the targeting ligand. In general, the compounds bound to the surface of the targeted delivery system will be ligands and receptors which will allow the targeted delivery system to find and "home in" on the desired cells. A ligand may be any compound of interest which will bind to another compound, such as a receptor.

15 The therapeutic agents useful in the method of the invention can be administered parenterally by injection or by gradual perfusion over time. Administration may be intravenously, intraperitoneally, intramuscularly, subcutaneously, intracavity, or transdermally.

Preparations for parenteral administration include sterile aqueous or non-  
20 aqueous solutions, suspensions, and emulsions. Examples of non-aqueous solvents are propylene glycol, polyethylene glycol, vegetable oils such as olive oil, and injectable organic esters such as ethyl oleate. Aqueous carriers include water, alcoholic/aqueous solutions, emulsions or suspensions, including saline and buffered media. Parenteral vehicles include sodium chloride solution, Ringer's dextrose,  
25 dextrose and sodium chloride, lactated Ringer's intravenous vehicles include fluid and nutrient replenishers, electrolyte replenishers (such as those based on Ringer's dextrose), and the like. Preservatives and other additives may also be present such as, for example, antimicrobials, anti-oxidants, chelating agents and inert gases and the like.

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The invention also includes a pharmaceutical composition comprising a therapeutically effective amount of T $\beta$ 4 or a T $\beta$ 4 isoform in a pharmaceutically acceptable carrier. Such carriers include those listed above with reference to parenteral administration.

- 5           The actual dosage or reagent, formulation or composition that modulates a tissue repair process, fibrotic disorder, a sclerotic disorder, a cell proliferative disorder, or wound healing depends on many factors, including the size and health of a subject. However, one of ordinary skill in the art can use the following teachings describing the methods and techniques for determining clinical dosages (Spilker B.,
- 10 *Guide to Clinical Studies and Developing Protocols*, Raven Press Books, Ltd., New York, 1984, pp. 7-13, 54-60; Spilker B., *Guide to Clinical Trials*, Raven Press, Ltd., New York, 1991, pp. 93-101; Craig C., and R. Stitzel, eds., *Modern Pharmacology*, 2d ed., Little, Brown and Co., Boston, 1986, pp. 127-33; T. Speight, ed., *Avery's Drug Treatment: Principles and Practice of Clinical Pharmacology and Therapeutics*, 3d
- 15 ed., Williams and Wilkins, Baltimore, 1987, pp. 50-56; R. Tallarida, R. Raffa and P. McGonigle, *Principles in General Pharmacology*, Springer-Verlag, New York, 1988, pp. 18-20) or to determine the appropriate dosage to use.

#### **Antibodies that Bind to T $\beta$ 4**

- Antibodies to T $\beta$ 4 peptide or fragments could be valuable as diagnostic
- 20 tools to aid in the detection of diseases in which T $\beta$ 4 is a pathological factor. Further, use of antibodies which bind to T $\beta$ 4 and inhibit or prevent the actions of T $\beta$ 4 are included in the present invention. Therapeutically, antibodies or fragments of the antibody molecule could also be used to neutralize the biological activity of T $\beta$ 4 in diseases where T $\beta$ 4 is over expressed. Such antibodies can recognize an epitope of
- 25 T $\beta$ 4 or fragments thereof suitable for antibody recognition and neutralization of T $\beta$ 4 activity. As used in this invention, the term "epitope" refers to an antigenic determinant on an antigen, such as a T $\beta$ 4 peptide, to which the paratope of an antibody, such as a T $\beta$ 4-specific antibody, binds. Antigenic determinants usually consist of chemically active surface groupings of molecules, such as amino acids or

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sugar side chains, and can have specific three-dimensional structural characteristics, as well as specific charge characteristics.

Preparation of an antibody requires a substantially purified moiety that can provide an antigenic determinant. The term "substantially pure" as used herein refers to T $\beta$ 4, or variants thereof, which is substantially free of other proteins, lipids, carbohydrates or other materials with which it is naturally associated. Substantially purified or "isolated" refers to molecules, either nucleic or amino acid sequences, that are removed from their natural environment, isolated or separated, and are at least 60% free, preferably 75% free, and most preferably 90% free from other components with which they are naturally associated. One skilled in the art can isolate T $\beta$ 4 or a T $\beta$ 4 isoform using standard techniques for protein purification. The substantially pure peptide will yield a single major band on a non-reducing polyacrylamide gel. The purity of the T $\beta$ 4 peptide can also be determined by amino-terminal amino acid sequence analysis. T $\beta$ 4 or a T $\beta$ 4 isoform peptide includes functional fragments of the peptide, as long as the activity of T $\beta$ 4 or a T $\beta$ 4 isoform remains. Smaller peptides containing the biological activity of T $\beta$ 4 or a T $\beta$ 4 isoform are included in the invention. As used in the present invention, the term "antibody" includes, in addition to conventional antibodies, such protein fragments that have the ability to recognize specifically and bind the T $\beta$ 4 protein or variants thereof. Regions of the gene that differ at the protein level are well defined. A protein can be raised by expression of the wild type (wt) gene or of the variants, or, preferably, fractions thereof. For example, the nucleic acid sequence can be cloned into expression vectors. According to this embodiment, the sequence of interest can first be obtained by employing PCR, as described above, or from a synthetic gene construction with overlapping and ligated synthetic oligonucleotides. Another alternative would involve synthesis of a short peptide. All those methodologies are well known to one skilled in the art. See, for example, Ausubel *et al.*, CURRENT PROTOCOLS IN MOLECULAR BIOLOGY, Volumes 1 and 2 (1987), with supplements, and Maniatis *et al.*, MOLECULAR CLONING, A LABORATORY MANUAL, Cold Spring Harbor Laboratory, all of which are incorporated herein by reference.

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The invention provides a method for detecting T $\beta$ 4, or variants thereof, which includes contacting an anti-T $\beta$ 4 antibody with a sample suspected of containing T $\beta$ 4, (*e.g.*, cell or protein) and detecting binding to the antibody. An antibody which binds to T $\beta$ 4 peptide is labeled with a compound which allows  
5 detection of binding to T $\beta$ 4. There are many different labels and methods of labeling known to those of ordinary skill in the art. Examples of the types of labels which can be used in the present invention include enzymes, radioisotopes, fluorescent compounds, colloidal metals, chemiluminescent compounds, phosphorescent compounds, and bioluminescent compounds. Those of ordinary skill in the art will  
10 know of other suitable labels for binding to the antibody, or will be able to ascertain such, using routine experimentation. For purposes of the invention, an antibody specific for T $\beta$ 4 peptide may be used to detect the level of T $\beta$ 4 in biological fluids and tissues. Any specimen containing a detectable amount of antigen can be used. The level of T $\beta$ 4 in the suspect cell can be compared with the level in a normal cell to  
15 determine whether the subject is predisposed to a T $\beta$ 4 associated increase in angiogenesis or wound healing.

Use of antibodies for the diagnostic methods of the invention includes, for example, immunoassays in which the antibodies can be utilized in liquid phase or bound to a solid phase carrier. In addition, the antibodies in these immunoassays can  
20 be detectably labeled in various ways. Examples of types of immunoassays which can utilize antibodies of the invention are competitive and non-competitive immunoassays in either a direct or indirect format. Examples of such immunoassays are the radioimmunoassay (RIA) and the sandwich (immunometric) assay. Detection of the antigens using the antibodies of the invention can be done utilizing  
25 immunoassays which are run in either the forward, reverse, or simultaneous modes, including immunohistochemical assays on physiological samples. Those of skill in the art will know, or can readily discern, other immunoassay formats without undue experimentation.

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T $\beta$ 4 antibodies can be bound to many different carriers and used to detect the presence of an antigen comprising the peptide of the invention. Examples of well-known carriers include glass, polystyrene, polypropylene, polyethylene, dextran, nylon, amylases, natural and modified celluloses, polyacrylamides, agaroses and  
5 magnetite. The nature of the carrier can be either soluble or insoluble for purposes of the invention. Those skilled in the art will know of other suitable carriers for binding antibodies, or will be able to ascertain such, using routine experimentation.

Another technique which may also result in greater sensitivity consists of coupling the antibodies to low molecular weight haptens. These haptens can then be  
10 specifically detected by means of a second reaction. For example, it is common to use such haptens as biotin, which reacts with avidin, or dinitrophenyl, puridoxal, and fluorescein, which can react with specific antihapten antibodies.

The invention includes use of antibodies immunoreactive with T $\beta$ 4 peptide or functional fragments thereof. Antibody which consists essentially of pooled  
15 monoclonal antibodies with different epitopic specificities, as well as distinct monoclonal antibody preparations are provided. Monoclonal antibodies are made from antigen containing fragments of the protein by methods well known to those skilled in the art (Kohler, *et al.*, *Nature*, 256:495, 1975). The term antibody as used in this invention is meant to include intact molecules as well as fragments thereof, such  
20 as Fab and F(ab')<sub>2</sub>, Fv and SCA fragments which are capable of binding an epitopic determinant on T $\beta$ 4.

(1) An Fab fragment consists of a monovalent antigen-binding fragment of an antibody molecule, and can be produced by digestion of a whole antibody molecule with the enzyme papain, to yield a fragment consisting of an intact light  
25 chain and a portion of a heavy chain.

(2) An Fab' fragment of an antibody molecule can be obtained by treating a whole antibody molecule with pepsin, followed by reduction, to yield a molecule consisting of an intact light chain and a portion of a heavy chain. Two Fab' fragments are obtained per antibody molecule treated in this manner.

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(3) An (Fab')<sub>2</sub> fragment of an antibody can be obtained by treating a whole antibody molecule with the enzyme pepsin, without subsequent reduction. A (Fab')<sub>2</sub> fragment is a dimer of two Fab' fragments, held together by two disulfide bonds.

(4) An Fv fragment is defined as a genetically engineered fragment  
5 containing the variable region of a light chain and the variable region of a heavy chain expressed as two chains.

(5) A single chain antibody ("SCA") is a genetically engineered single chain molecule containing the variable region of a light chain and the variable region of a heavy chain, linked by a suitable, flexible polypeptide linker.

10 Alternatively, a therapeutically or diagnostically useful anti-Tβ4 antibody may be derived from a "humanized" monoclonal antibody. Humanized monoclonal antibodies are produced by transferring mouse complementary determining regions from heavy and light variable chains of the mouse immunoglobulin into a human variable domain, and then substituting human residues in the framework regions of  
15 the murine counterparts. The use of antibody components derived from humanized monoclonal antibodies obviates potential problems associated with the immunogenicity of murine constant regions. General techniques for cloning murine immunoglobulin variable domains are described, for example, by Orlandi *et al.*, *Proc. Natl. Acad. Sci. USA* 86: 3833 (1989), which is hereby incorporated in its entirety by  
20 reference. Techniques for producing humanized monoclonal antibodies are described, for example, by Jones *et al.*, *Nature* 321: 522 (1986); Riechmann *et al.*, *Nature* 332: 323 (1988); Verhoeven *et al.*, *Science* 239: 1534 (1988); Carter *et al.*, *Proc. Nat'l Acad. Sci. USA* 89: 4285 (1992); Sandhu, *Crit. Rev. Biotech.* 12: 437 (1992); and Singer *et al.*, *J. Immunol.* 150: 2844 (1993), which are hereby incorporated by  
25 reference.

Antibodies of the invention also may be derived from human antibody fragments isolated from a combinatorial immunoglobulin library. See, for example, Barbas *et al.*, METHODS: A COMPANION TO METHODS IN ENZYMOLOGY, VOL. 2, page 119 (1991); Winter *et al.*, *Ann. Rev. Immunol.* 12: 433 (1994), which  
30 are hereby incorporated by reference. Cloning and expression vectors that are useful

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for producing a human immunoglobulin phage library can be obtained, for example, from STRATAGENE Cloning Systems (La Jolla, CA).

**Methods and Compositions for Treating or Diagnosing T $\beta$ 4-Associated Disorders**

In another embodiment of the invention, a method of diagnosing a  
5 pathological state in a subject suspected of having a pathology characterized by a disorder associated with T $\beta$ 4 is provided. The method includes obtaining a sample suspected of containing T $\beta$ 4 from the subject, determining the level of T $\beta$ 4 in the sample and comparing the level of T $\beta$ 4 in the sample to the level of T $\beta$ 4 in a normal standard sample. Such conditions include, but are not limited to subjects having cell  
10 proliferative disorders, recurrent wounds, tissue repair disorders, fibrotic tissue disorders, chronic ulcers and other disorders described herein. Such disorders further include those associated with the various T $\beta$ 4 isoforms, known or not yet identified.

The term "cell-proliferative disorder" denotes malignant as well as non-malignant cell populations which often appear to differ from the surrounding tissue  
15 both morphologically and genotypically. Malignant cells (*i.e.* cancer) develop as a result of a multistep process. Such disorders may be detected using the methods of the current invention. For example, a sample suspected of containing T $\beta$ 4 is obtained from a subject, the level of T $\beta$ 4 peptide is determined and compared with the level of T $\beta$ 4 peptide in a normal tissue sample. The level of T $\beta$ 4 can be determined by any  
20 number of methods including, for example, immunoassay using anti-T $\beta$ 4 peptide antibodies. Other variations of such assays include radioimmunoassay (RIA), ELISA and immunofluorescence. Alternatively, nucleic acid probes can be used to detect and quantify T $\beta$ 4 peptide mRNA for the same purpose. Such detection methods are standard in the art.

25 In another embodiment, the invention provides a method for ameliorating a wound healing disorder associated with T $\beta$ 4 or a T $\beta$ 4 isoform, including treating a subject having the disorder with a composition that regulates T $\beta$ 4 activity. The term "ameliorate" denotes a lessening of the detrimental effect of the disease-inducing response in the subject receiving therapy. Where the disease is due to an abnormally



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high level of T $\beta$ 4, the administration of an agent, such as an antagonist of T $\beta$ 4 activity, may be effective in decreasing the amount of T $\beta$ 4 activity. Alternatively, where the disease is due to an abnormally low level of T $\beta$ 4, the administration of T $\beta$ 4 or an agent that increases T $\beta$ 4 activity, such as an agonist, may be effective in  
5 increasing the amount of T $\beta$ 4 activity.

In yet another embodiment, the invention provides a method of treating a subject having a wound healing disorder characterized by recurrent or slow to heal wounds or wounds that are chronic non-healing wounds associated with altered T $\beta$ 4 or T $\beta$ 4 isoform gene expression in a subject. The method includes administering to a  
10 subject having the disorder a wound-healing effective amount of an agent which modulates T $\beta$ 4 gene expression, thereby treating the disorder. The term "modulate" refers to inhibition or suppression of T $\beta$ 4 expression when T $\beta$ 4 is over expressed, and induction of expression when T $\beta$ 4 is under expressed. The term "wound-healing effective amount" means that amount of T $\beta$ 4 agent which is effective in modulating  
15 T $\beta$ 4 gene expression resulting in reducing the symptoms of the T $\beta$ 4 associated wound healing disorder.

An agent which modulates T $\beta$ 4 or T $\beta$ 4 isoform gene expression may be a polynucleotide for example. The polynucleotide may be an antisense, a triplex agent, or a ribozyme. For example, an antisense may be directed to the structural gene  
20 region or to the promoter region of T $\beta$ 4 may be utilized.

When a wound healing disorder is associated with the expression of T $\beta$ 4, a therapeutic approach which directly interferes with the translation of T $\beta$ 4 mRNA into protein is possible. For example, an antisense nucleic acid or a ribozyme can be used to bind to the T $\beta$ 4 RNA or to cleave it. Antisense RNA or DNA molecules bind  
25 specifically with a targeted gene's RNA message, interrupting the expression of that gene's protein product. The antisense binds to the mRNA forming a double stranded molecule which cannot be translated by the cell. Antisense oligonucleotides of about 15-25 nucleotides are preferred since they are easily synthesized and have an inhibitory effect just like antisense RNA molecules. In addition, chemically reactive  
30 group, such as iron-linked ethylenediaminetetraacetic acid (EDTA-Fe) can be

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attached to an antisense oligonucleotide, causing cleavage of the RNA at the site of hybridization. These and other uses of antisense methods to inhibit the *in vitro* translation of genes are well known in the art (Marcus-Sakura, *Anal., Biochem.*, 172:289, 1988).

5           Antisense nucleic acids are DNA or RNA molecules that are complementary to at least a portion of a specific mRNA molecule (Weintraub, *Scientific American*, 262:40, 1990). In the cell, the antisense nucleic acids hybridize to the corresponding mRNA, forming a double-stranded molecule. The antisense nucleic acids interfere with the translation of the mRNA, since the cell will not  
10   translate a mRNA that is double-stranded. Antisense oligomers of about 15 nucleotides are preferred, since they are easily synthesized and are less likely to cause problems than larger molecules when introduced into the target T $\beta$ 4 producing cell. The use of antisense methods to inhibit the *in vitro* translation of genes is well known in the art (Marcus-Sakura, *Anal. Biochem.*, 172:289, 1988).

15           Use of an oligonucleotide to stall transcription is known as the triplex strategy since the oligomer winds around double-helical DNA, forming a three-strand helix. Therefore, these triplex compounds can be designed to recognize a unique site on a chosen gene (Maher, *et al.*, *Antisense Res. and Dev.*, 1(3):227, 1991; Helene, C., *Anticancer Drug Design*, 6(6):569, 1991).

20           Ribozymes are RNA molecules possessing the ability to specifically cleave other single-stranded RNA in a manner analogous to DNA restriction endonucleases. Through the modification of nucleotide sequences which encode these RNAs, it is possible to engineer molecules that recognize specific nucleotide sequences in an RNA molecule and cleave it (Cech, *J. Amer. Med. Assn.*, 260:3030,  
25   1988). A major advantage of this approach is that, because they are sequence-specific, only mRNAs with particular sequences are inactivated.

          There are two basic types of ribozymes namely, *tetrahymena*-type (Hasselhoff, *Nature*, 334:585, 1988) and "hammerhead"-type. *Tetrahymena*-type ribozymes recognize sequences which are four bases in length, while "hammerhead"-  
30   type ribozymes recognize base sequences 11-18 bases in length. The longer the

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recognition sequence, the greater the likelihood that the sequence will occur exclusively in the target mRNA species.

These and other uses of antisense methods to inhibit the *in vivo* translation of genes are well known in the art (*e.g.*, De Mesmaeker, *et al.*, 1995. Backbone modifications in oligonucleotides and peptide nucleic acid systems. *Curr. Opin. Struct. Biol.* 5:343-355; Gewirtz, A.M., *et al.*, 1996b. Facilitating delivery of antisense oligodeoxynucleotides: Helping antisense deliver on its promise; *Proc. Natl. Acad. Sci. U.S.A.* 93:3161-3163; Stein, C.A. A discussion of G-tetrads 1996. Exploiting the potential of antisense: beyond phosphorothioate oligodeoxynucleotides. *Chem. and Biol.* 3:319-323).

Delivery of antisense, triplex agents, ribozymes, competitive inhibitors and the like can be achieved using a recombinant expression vector such as a chimeric virus or a colloidal dispersion system. Various viral vectors which can be utilized for gene therapy as taught herein include adenovirus, herpes virus, vaccinia, or, preferably, an RNA virus such as a retrovirus. Preferably, the retroviral vector is a derivative of a murine or avian retrovirus. Examples of retroviral vectors in which a single foreign gene can be inserted include, but are not limited to: Moloney murine leukemia virus (MoMuLV), Harvey murine sarcoma virus (HaMuSV), murine mammary tumor virus (MuMTV), and Rous Sarcoma Virus (RSV). A number of additional retroviral vectors can incorporate multiple genes. All of these vectors can transfer or incorporate a gene for a selectable marker so that transduced cells can be identified and generated. By inserting a polynucleotide sequence of interest into the viral vector, along with another gene which encodes the ligand for a receptor on a specific target cell, for example, the vector is now target specific. Retroviral vectors can be made target specific by inserting, for example, a polynucleotide encoding a sugar, a glycolipid, or a protein. Preferred targeting is accomplished by using an antibody to target the retroviral vector. Those of skill in the art will know of, or can readily ascertain without undue experimentation, specific polynucleotide sequences which can be inserted into the retroviral genome to allow target specific delivery of the retroviral vector containing the antisense polynucleotide.

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Since recombinant retroviruses are defective, they require assistance in order to produce infectious vector particles. This assistance can be provided, for example, by using helper cell lines that contain plasmids encoding all of the structural genes of the retrovirus under the control of regulatory sequences within the LTR.

- 5 These plasmids are missing a nucleotide sequence which enables the packaging mechanism to recognize an RNA transcript for encapsidation. Helper cell lines which have deletions of the packaging signal include but are not limited to  $\Psi$ 2, PA317 and PA12, for example. These cell lines produce empty virions, since no genome is packaged. If a retroviral vector is introduced into such cells in which the packaging  
10 signal is intact, but the structural genes are replaced by other genes of interest, the vector can be packaged and vector virion produced.

- Alternatively, NIH 3T3 or other tissue culture cells can be directly transfected with plasmids encoding the retroviral structural genes *gag*, *pol* and *env*, by conventional calcium phosphate transfection. These cells are then transfected with  
15 the vector plasmid containing the genes of interest. The resulting cells release the retroviral vector into the culture medium.

- A targeted delivery system for delivery of nucleic acids as described herein includes a colloidal dispersion system. Colloidal dispersion systems include macromolecule complexes, nanocapsules, microspheres, beads, gene activated  
20 matrices and lipid-based systems including oil-in-water emulsions, micelles, mixed micelles, and liposomes. The preferred colloidal system of this invention is a liposome. Liposomes are artificial membrane vesicles which are useful as delivery vehicles *in vitro* and *in vivo*. It has been shown that large unilamellar vesicles (LUV), which range in size from 0.2-4.0  $\mu$ m can encapsulate a substantial percentage of an  
25 aqueous buffer containing large macromolecules. RNA, DNA and intact virions can be encapsulated within the aqueous interior and be delivered to cells in a biologically active form (Fraley, *et al.*, *Trends Biochem. Sci.*, 6:77, 1981). In addition to mammalian cells, liposomes have been used for delivery of polynucleotides in plant, yeast and bacterial cells. In order for a liposome to be an efficient gene transfer  
30 vehicle, the following characteristics should be present: (1) encapsulation of the genes

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of interest at high efficiency while not compromising their biological activity; (2) preferential and substantial binding to a target cell in comparison to non-target cells; (3) delivery of the aqueous contents of the vesicle to the target cell cytoplasm at high efficiency; and (4) accurate and effective expression of genetic information

5 (Mannino, *et al.*, *Biotechniques*, 6:682, 1988).

Pathologically, T $\beta$ 4 may be involved in diseases in which there is an overgrowth of blood vessels, such as cancer, tumor formation and growth, diabetic retinopathy, neovascular glaucoma, rheumatoid arthritis and psoriasis.

The ingrowth of capillaries and ancillary blood vessels is essential for  
10 growth of solid tumors and is thus an unwanted physiological response which facilitates the spread of malignant tissue and metastases. Inhibition of angiogenesis and the resultant growth of capillaries and blood vessels is therefore a component of effective treatment of malignancy in use of treatment of cancer patients.

Thus, in another embodiment, the invention provides a method of  
15 inhibiting angiogenesis in a subject, including administering to the subject a composition containing an agent which regulates T $\beta$ 4 activity. The composition may include agents that regulate angiogenesis, for example agents that affect thymosin  $\alpha$ 1, PDGF, VEGF, IGF, FGF and TGF $\beta$ . For example, the inhibition of angiogenesis and endothelial cell migration can be beneficial in controlling the growth of solid tumors.  
20 Most, if not all solid tumors, like normal tissue, require a steady and sufficient blood supply for optimal growth. Tumors are known to make use of angiogenic growth factors to attract new blood vessels and ascertain supply with sufficient amounts of nutrients to sustain their growth. Many tumors are well vascularized and the inhibition of the formation of an adequate blood supply to the tumor by inhibition of  
25 tumor vascularization, as a result of inhibition of angiogenesis, is beneficial in tumor growth control. Without a strong blood supply, rapid and prolonged growth of tumor tissue cannot be sustained. Thus, agents that inhibit T $\beta$ 4 activity may be used to prevent neoplastic growth. The T $\beta$ 4 inhibiting agent may be administered orally, parenterally, topically, intravenously, or systemically. In addition, for inhibiting  
30 tumor cell proliferation and tumor growth, the agent may be administered locally

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directly to the tumor or as a part of a deposited slow release formulation.

Administration may be on a daily basis for as long as needed to inhibit angiogenesis, endothelial cell proliferation, tumor cell proliferation or tumor growth. Alternatively, a slow release formulation may continue for as long as needed to control tumor

5 growth. This dosage regimen may be adjusted to provide the optimum therapeutic response. For example, several divided doses may be administered daily or the dose may be proportionally reduced as indicated by the exigencies of the therapeutic situation.

In this regard, the compositions of this invention that are useful as  
10 inhibitors of angiogenesis, endothelial cell proliferation, tumor cell proliferation and tumor growth contain a pharmaceutically acceptable carrier and an amount of T $\beta$ 4 modulating agent effective to inhibit tumor or endothelial cell proliferation. Such compositions may also include preservatives, antioxidants, immunosuppressants and other biologically and pharmaceutically effective agents which do have effects on  
15 tumor growth but which do not exert a detrimental effect on the T $\beta$ 4 modulating agent. For treatment of tumor cells the composition may include a chemotherapeutic agent, for example an anti-cancer agent which selectively kills the faster replicating tumor cells, many of which are known and clinically used. Exemplary anticancer agents include mephalan, cyclophosphamide, methotrexate, adriamycin and  
20 bleomycin.

**Screen for compounds which modulate T $\beta$ 4 activity**

In another embodiment, the invention provides a method for identifying a compound that modulates T $\beta$ 4 activity, angiogenesis activity or wound healing activity. The method includes incubating components including the compound and  
25 T $\beta$ 4 under conditions sufficient to allow the components to interact and determining the effect of the compound on T $\beta$ 4 activity before and after incubating in the presence of the compound. Compounds that affect T $\beta$ 4 activity (e.g., antagonists and agonists) include peptides, peptidomimetics, polypeptides, chemical compounds, minerals such

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as zincs, and biological agents. T $\beta$ 4 activity can be assayed using the methodology as described in the present Examples.

The present Examples are meant to illustrate, but not limit the scope of the appended claims. Accordingly, one skilled in the art will recognize a number of  
5 equivalent materials and methods, which are intend to be covered by the present invention and disclosure.

### EXAMPLE 1

#### In vivo wound healing is accelerated by T $\beta$ 4

T $\beta$ 4, whether administered topically or intraperitoneal, significantly  
10 accelerated wound healing as compared to untreated wounds (Fig. 2 and 3). Full thickness 8 mm punch biopsy wounds were made on the dorsal surface of rats as previously reported (Bhartiya *et al.*, *J. Cell. Physiol.* 150:312, 1992; Sihhu *et al.*, *J. Cell. Physiol.* 169:108, 1996) and T $\beta$ 4 was given topically at the time of wounding (5  $\mu$ g in 50  $\mu$ l) and again after 48 hours. Controls for the topical treatment received  
15 identical amounts of saline at the time of wounding and at 48 hours. Additional rats received intraperitoneal injections at the time of wounding (60  $\mu$ g in 300  $\mu$ l) and again every other day (*e.g.*, days 0, 2, 4, and 6). Controls for these animals received identical amounts of saline intra-peritoneally on the same injection schedule. On days 4 and 7 post-wounding, measurements were made on the wound size. At days 8 and 9  
20 post-wounding, tissue was collected and fixed in 10% buffered formalin. The samples were sectioned and stained with H&E and Masson's Trichrome (American Histolabs, Gaithersburg, MD).

Histological sections were used to measure the re-epithelialization and the contraction of the wound using an ocular micrometer. Epidermal migration was  
25 determined by measuring the lengths of the tongues of epithelium migrating from either side of the wound over the wound bed from the zone of proliferation at the margin of the uninjured and wounded skin. Epidermal thickness was also measured beginning at the junction of the uninjured and proliferating epidermis. The thickness was measured vertically from the basement membrane to the most superficial layer of

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the migrating epidermis at every 200 microns. The mean epidermal thickness of each migrating tongue of epidermis was then computed from each wound. Vessel counts were performed by first identifying vascular spaces by their endothelial lining. All such vessels in the wound bed were counted including those at the junction of the dermis and the subcutis, since angiogenesis into the wounds occurs to a great extent from these vessels. The numbers were averaged into vessel counts per 10 high powered fields (40x).

The effect of T $\beta$ 4 on wound healing was studied in a full thickness cutaneous rat wound model. FIG. 1 shows a diagram of the wound site that extends from the epidermis to the fat/muscle layer. This model allowed measurement of two parameters: the re-epithelialization (gap) and the contraction (width) of the wound. Wounds treated topically with T $\beta$ 4 showed about a 15% decrease in width and about 15% decrease in gap in the treated versus controls (FIG. 2 and 3, respectively).

Figure 2 shows a 15% decrease in wound width as compared to the saline controls as early as 4 days after wounding and continued until day 7. Intraperitoneal injection of T $\beta$ 4 resulted in a 18% decrease in wound width relative to saline treated controls at day 4 and 11% decrease at day 7. This trend was observed on the 4th day post wounding and continued through day 7 (\*P $\leq$ 0.0001, \*\*P $\leq$ 0.08, significant difference from media alone, student's t-test). These data demonstrate that T $\beta$ 4, when given either topically or systemically, increases wound re-epithelialization and contraction. Both topical and systemic treatment are equally effective. Lower doses of T $\beta$ 4 were tested including 2.5  $\mu$ g and 0.5  $\mu$ g in 50  $\mu$ l for topical and 30  $\mu$ g and 6  $\mu$ g in 300  $\mu$ l for intraperitoneal injection but reduced or no effect, respectively, was observed on wound healing.

Figure 3 shows an 18% decrease in gap length as compared to saline controls when T $\beta$ 4 is administered topically, as early as 4 days after wounding. This trend continued to termination at day 7 (\*P $\leq$ 0.04, student's t-test). Intraperitoneal injections resulted in a 42% decrease in gap size relative to saline treated controls. This decrease was observed on the 4th day post wounding and continued through day 7 (\*\*P $\leq$ 0.0007, student's t-test). The increase in re-epithelialization was observed in



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wounds treated for 7 days and the rate of gap closure was slightly accelerated over that observed at day 4. A 62% decrease in gap size was observed in the T $\beta$ 4-treated wounds. Quantitation of epidermal migration showed a statistically significant 1.5 fold increase in migration of epidermal tongues over the wound bed after topical treatment (Table 1). Quantitation of epithelial migration in intraperitoneally treated wounds showed a statistically significant increase in migration of epidermal tongues as compared to controls (Table 1). There was no difference in the thickness of the migrating epidermis between either of the T $\beta$ 4 treatments and the control (Table 1). Histological sections of the wounds clearly show increased re-epithelialization in the treated wounds as compared to controls in 7 day wounds (FIG 4).

Table 1: Morphometric Measurements of Control and Thymosin  $\beta$ 4 Treated Samples

Parameter	Control	I.P.	Topical
Epidermal Migration ( $\mu$ m)	2403.3 $\pm$ 9.7	3168.3 $\pm$ 38.4*	3668.7 $\pm$ 56.6*
Epidermal Thickness ( $\mu$ m)	128.2 $\pm$ 19.3	135.0 $\pm$ 11.7	142.3 $\pm$ 19.8
Vessels/10 HPF	1364.0 $\pm$ 15.0	2415.0 $\pm$ 24.3*	2186.0 $\pm$ 11.8*

HPF: high power field. \*P $\leq$ 0.00001 by Welch's t-test, significantly different than control.

FIG. 4 shows a comparison of typical control (D) and T $\beta$ 4-treated (E and F) sections of 7 day wounds. Treatment with T $\beta$ 4 resulted in considerable capillary ingrowth (FIG 4E and F, arrows). Vessel counts showed a significant (about 2 fold) increase in the number of vessels in T $\beta$ 4 treated wounds (Table 1). No increases in the number of macrophages in the wounds were observed. There was no apparent increase in the accumulation/biosynthesis of collagen in treated -T $\beta$ 4 wounds (Fig. 5B and C vs A) supporting a decreased wound width and supporting a role for T $\beta$ 4 in wound contraction. Both the topical and systemically treated wound appeared similar although the wound contraction proceeded slightly more quickly with the topical treatment.

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Reduction of the wound size was observed in both experimental groups as compared to control groups (Fig. 2 - 4). More and larger blood vessels were noted in the experimental groups as compared to the controls (Fig. 4). Additionally, an increase in the accumulation/biosynthesis of collagen by T $\beta$ 4 treated wounds as compared to control suggests a role for T $\beta$ 4 in wound contraction and extracellular matrix deposition. Histological staining of these wounds demonstrated an increase in collagen density and extracellular matrix deposition when compared to controls. (Fig. 5).

## EXAMPLE 2

### 10        Migration Assays of Keratinocytes

Primary keratinocytes were prepared from either Balb/c or CD-1 newborn mice as described previously (Dlugosz *et al.*, 1995). Cells were plated in calcium- and magnesium-free Eagle's Minimal Essential Medium (EMEM) containing 8% fetal calf serum treated with 8% Chelex (Bio-Rad Laboratories, Hercules, CA), 20 units/ml penicillin-streptomycin, and the calcium concentration was adjusted to 0.25 mM. The following day, cultures were washed with calcium- and magnesium-free phosphate buffered saline, treated briefly with Trypsin (Life Technologies, Gaithersburg, MD), washed with culture medium and resuspended in EMEM containing 0.05 mM calcium. Cells were used immediately in migration assays.

20        Keratinocyte migration assays were carried out in Boyden chamber using 12  $\mu$ m pore polyester membranes (Poretics, Livermore, CA) coated with a 0.1 mg/ml solution of collagen IV in dH<sub>2</sub>O (Trevigen, Gaithersburg, MD). Filters were then dried at least 1 h. Cells were harvested using Versene or Trypsin (Life Technologies, Gaithersburg, MD) and resuspended in Eagle's minimal essential medium with 0.05 mM Ca<sup>2+</sup>. The bottom chamber was loaded with EMEM containing 0.01, 0.1, 10, 100, and 1000 ng/ml of synthetic T $\beta$ 4. Conditioned medium from primary dermal fibroblasts and/or keratinocyte growth factor was added to several wells as a positive control. Cells were added to the upper chamber at a concentration of 50,000 cells per well. Chambers were incubated at 35 C/7% CO<sub>2</sub> for 4-5 hours and the filters were

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then fixed and stained using Diff-Quik (Baxter Healthcare Corporation, McGraw Park, IL). The cells that migrated through the filter were quantitated by counting the center of each well at 10x using an Olympus CK2 microscope. Each condition was assayed in triplicate wells and each experiment was repeated four times with different  
5 preparations of cells.

The results demonstrated that keratinocyte migrated in response to T $\beta$ 4 after 4-5 hours of exposure. Migration was enhanced 2-3 fold ( $P \leq 0.003$ ) over migration in the presence of media alone (FIG. 6) and at the maximal responding dose exceeded the positive control. The effect of T $\beta$ 4 on migration, while showing slightly  
10 different dose curves depending on the cell preparation and source, clearly showed a biphasic pattern with 1000 ng/ml and 0.01 ng/ml showing the most migration and the middle doses showing less stimulation (but still greater than control media) in all 4 assays.

15

### EXAMPLE 3

#### Migration Assays of Corneal Epithelial Cells

Corneal Epithelial Cell migration assays were carried out in Boyden chamber using 12  $\mu$ m pore polyester membranes (Poretics, Livermore, CA) coated with a 0.1 mg/ml solution of collagen IV in dH<sub>2</sub>O (Trevigen, Gaithersburg, MD).  
20 Filters were then dried at least 1 h. Cells were cultured and resuspended in Eagle's Minimal Essential Medium with 0.05 mM Ca<sup>2+</sup>. The bottom chamber was loaded with EMEM containing 0.01, 0.1, 10, 100, and 1000 ng/ml of synthetic T $\beta$ 4. Conditioned medium from primary dermal fibroblasts and/or keratinocyte growth factor was added to several wells as a positive control. Cells were added to the upper  
25 chamber at a concentration of 50,000 cells per well. Chambers were incubated at 35 C/7% CO<sub>2</sub> for 4-5 hours and the filters were then fixed and stained using Diff-Quik (Baxter Healthcare Corporation, McGraw Park, IL). The cells that migrated through the filter were quantitated by counting the center of each well at 10x using an Olympus CK2 microscope. Each condition was assayed in triplicate wells and each  
30 experiment was repeated four times with different preparations of cells.

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The results demonstrated that corneal epithelial cell migrated in response to T $\beta$ 4 after 4-5 hours of exposure. Migration was enhanced 2-3 fold over migration in the presence of media alone (FIG. 7) with the highest level of migration seen at 100 ng/ml of T $\beta$ 4.

5 **EXAMPLE 4**

### In vivo Corneal Re-Epithelialization

To determine the effect of Tβ4 on corneal re-epithelialization *in vivo*, Rat corneas were de-epithelialized and treated with Tβ4. Filters were soaked in heptanol, applied to the eye for 30 seconds, and then the epithelium was scraped. Various concentration of Tβ4 in saline was applied to the eye and at 24 hours the rats were sacrificed. The eyes were fixed, sectioned and the degree of corneal epithelial migration (as measured in pixels) was determined using a microscope with an internal caliper by a masked observer. The results demonstrate that re-epithelialization of the cornea was increased 2-fold over untreated control in the presence of about 1 to 25μg of Tβ4 (FIG. 8 and 9). In addition, it was noted that Tβ4 treated eyes had reduced inflammation compared to the non-treated corneas.

### EXAMPLE 5

### Impaired Healing Model

Thymosin  $\beta 4$  also enhanced wound healing in an impaired model. Steroid treatment reduces the rate of wound repair in mammals. Rats treated with steroids such as hydrocortisone serve as a model of impaired wound healing due to the delay observed in wound closure. Animals were injected intramuscularly everyday with hydrocortisone. Steroid treated rats showed a significant increase in the level of healing when T $\beta 4$  was added topically or injected intraperitoneally. At the initial time point, day 4, topically treated animals showed little response ( $\leq 7\%$  gap or width closure, N=3) compared to saline treatment. Intraperitoneal injection, however, resulted in a 28% decrease in gap size and a 14% decrease in wound width. At day 7, a response was observed with both topical treatment and intraperitoneal injection.

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The gap in topically treated animals decreased by 39% compared to saline treatment. The wound width decreased by 23%. Intraperitoneal injection resulted in a 26% decrease in gap size and a 10% decrease in wound width. Taken together, these demonstrate that T $\beta$ 4 is useful to treat chronic, as well as, acute wounds.

- 5           A number of embodiments of the present invention have been described. Nevertheless, it will be understood that various modifications may be made without departing from the spirit and scope of the invention. Accordingly, other embodiments are within the scope of the following claims.

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**WHAT IS CLAIMED IS:**

1. A method for promoting wound healing in a subject in need of such treatment comprising administering to the subject a wound-healing effective amount of a composition containing a wound healing polypeptide comprising the amino acid  
5 sequence LKKTET and conservative variants thereof having wound healing activity.
2. The method of claim 1, wherein the wound healing polypeptide is thymosin  $\beta$ 4 or an isoform of thymosin  $\beta$ 4.
3. The method of claim 2, wherein the composition further contains an agent that  
10 stimulates the production of thymosin  $\beta$ 4 peptide.
4. The method of claim 3, wherein the agent is transforming growth factor beta (TGF- $\beta$ ).
5. The method of claim 1, wherein the wound healing polypeptide is delivered systemically.
- 15 6. The method of claim 1, wherein the wound healing polypeptide is delivered topically.
7. The method of claim 6, wherein the wound healing polypeptide is contained in a topical formulation selected from the group consisting of a gel, cream, paste, lotion, spray, suspension, dispersion, salve, hydrogel and ointment.
- 20 8. The method of claim 1, wherein the wound healing polypeptide is recombinant or synthetic.

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9. The method of claim 2, wherein the isoform of thymosin  $\beta$ 4 is at least 70% homologous to thymosin  $\beta$ 4 peptide set forth as SEQ ID NO:1 in Figure 10.
10. The method of claim 9, wherein the isoform of thymosin  $\beta$ 4 is selected from the group consisting of: T $\beta$ 4<sup>ala</sup>, T $\beta$ 9, T $\beta$ 10, T $\beta$ 11, T $\beta$ 12, T $\beta$ 13, T $\beta$ 14 and T $\beta$ 15.
- 5 11. The method of claim 1, further comprising contacting the site of the wound with an agent which promotes wound healing.
12. The method of claim 11, wherein the agent is selected from the group consisting of IGF, IGF-1, IGF-2, IL-1, PDGF, FGF, KGF, VEGF, prothymosin  $\alpha$ , thymosin  $\alpha$ 1 or combinations thereof.
- 10 13. A method for promoting wound healing in a subject in need of such treatment comprising administering to the subject a wound-healing effective amount of a composition containing thymosin  $\beta$ 4 or an isoform of thymosin  $\beta$ 4.
14. The method of claim 13, wherein the composition further contains an agent that stimulates the production of thymosin  $\beta$ 4 peptide.
- 15 15. The method of claim 14, wherein the agent is transforming growth factor beta (TGF-b).
16. The method of claim 13, wherein the thymosin  $\beta$ 4 is delivered systemically.
17. The method of claim 13, wherein the thymosin  $\beta$ 4 is delivered topically.
18. The method of claim 17, wherein the thymosin  $\beta$ 4 is contained in a topical  
20 formulation selected from the group consisting of a gel, cream, paste, lotion, spray, suspension, dispersion, salve, hydrogel and ointment.

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19. The method of claim 13, wherein the thymosin  $\beta$ 4 is recombinant or synthetic.
20. The method of claim 13, wherein the isoform of thymosin  $\beta$ 4 is at least 70% homologous to thymosin  $\beta$ 4 peptide set forth as SEQ ID NO:1 in Figure 10.
21. The method of claim 13, wherein the isoform of thymosin  $\beta$ 4 is selected from the group consisting of: T $\beta$ 4<sup>ala</sup>, T $\beta$ 9, T $\beta$ 10, T $\beta$ 11, T $\beta$ 12, T $\beta$ 13, T $\beta$ 14 and T $\beta$ 15.
22. The method of claim 13, further comprising contacting the site of the wound with an agent which promotes wound healing.
23. A method for promoting wound healing in a tissue comprising contacting the tissue with a therapeutically effective amount of a composition containing a wound healing polypeptide comprising the amino acid sequence LKKTET and conservative variants thereof having wound healing activity.
24. The method of claim 23, wherein the wound healing polypeptide is thymosin  $\beta$ 4 or an isoform of thymosin  $\beta$ 4.
25. The method of claim 23, wherein the contacting is *in vivo* in a subject.
26. The method of claim 23, wherein the contacting is *ex vivo*.
27. The method of claim 23, wherein the subject is a mammal.
28. The method of claim 27, wherein the mammal is human.
29. The method of claim 24, wherein the composition further contains an agent that stimulates the production of thymosin  $\beta$ 4 peptide.



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30. The method of claim 29, wherein the agent is transforming growth factor beta (TGF-b).
31. The method of claim 29, wherein the agent is a mineral.
32. The method of claim 29, wherein the mineral is zinc.
- 5 33. The method of claim 23, wherein the wound healing polypeptide is delivered topically.
34. The method of claim 23, wherein the wound healing polypeptide is contained in a topical formulation selected from the group consisting of a gel, cream, paste, lotion, spray, suspension, dispersion, salve, hydrogel and ointment.
- 10 35. The method of claim 23, wherein the wound healing polypeptide is delivered systemically.
36. The method of claim 23, further comprising contacting the site of the tissue with an agent which promotes wound healing.
37. The method of claim 36, wherein the agent is selected from the group consisting  
15 of IGF, IGF-1, IGF-2, PDGF, FGF, KGF, VEGF, prothymosin  $\alpha$ , thymosin  $\alpha$ 1 or combinations thereof.
38. The method of claim 23, wherein the tissue is selected from the group consisting of epidermal, eye, uro-genital, gastro-intestinal, cardiovascular, muscle, connective, and neural.
- 20 39. The method of claim 23, wherein the tissue is skin tissue.

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40. The method of claim 23, wherein the tissue is eye tissue.
41. A method of inhibiting wound healing in a subject, comprising administering to the subject a composition containing an agent which regulates thymosin  $\beta$ 4 activity.
- 5 42. The method of claim 41, wherein the agent is an antibody.
43. The method of claim 42, wherein the antibody is polyclonal.
44. The method of claim 42, wherein the antibody is monoclonal.
45. A method of diagnosing a pathological state in a subject suspected of having pathology characterized by a wound healing disorder associated with thymosin  $\beta$ 4, comprising:
- 10     obtaining a sample suspected of containing thymosin  $\beta$ 4 from the subject;  
      detecting a level of thymosin  $\beta$ 4 in the sample; and  
      comparing the level of thymosin  $\beta$ 4 in the sample to the level of thymosin  $\beta$ 4 in a normal standard sample.
- 15 46. The method of claim 45, wherein the pathology is selected from the group consisting of fibrotic disease, ischemia, atherosclerosis and cell proliferative disorders.
47. A method for ameliorating a wound healing disorder associated with thymosin  $\beta$ 4, comprising treating a subject having the disorder, at the site of the disorder,
- 20     with an agent which regulates thymosin  $\beta$ 4 or the activity of a thymosin  $\beta$ 4 isoform.

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- 48 The method of claim 47, wherein the thymosin  $\beta$ 4 regulating agent is an antagonist of thymosin  $\beta$ 4 peptide.
49. The method of claim 48, wherein the antagonist is an antibody which specifically binds to thymosin  $\beta$ 4 peptide.
- 5 50. A method for identifying a compound which modulates wound healing, angiogenesis or cell migration activity, comprising contacting thymosin  $\beta$ 4 or an isoform of thymosin  $\beta$ 4 with a compound suspected of having thymosin  $\beta$ 4 modulating activity and detecting an effect on thymosin  $\beta$ 4 or thymosin  $\beta$ 4 isoform activity.
- 10 51 The method of claim 50, wherein the compound is an agonist of thymosin  $\beta$ 4 activity.
52. The method of claim 50, wherein the compound is an antagonist of thymosin  $\beta$ 4 activity.
- 15 53 A method of promoting epithelial cell migration, comprising contacting an epithelial cell with a composition comprising thymosin  $\beta$ 4 or an isoform of thymosin  $\beta$ 4.
54. The method of claim 53, wherein the epithelial cell is a skin cell.
55. The method of claim 54, wherein the skin cell is a keratinocyte.
56. The method of claim 53, wherein the epithelial cell is a corneal epithelial cell.
- 20 57. The method of claim 53, wherein the contacting is *in vivo*.

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58. The method of claim 57, wherein the contacting is topical.
59. The method of claim 57, wherein the contacting is systemic.
60. The method of claim 53, wherein the contacting is *in vitro* or ex vivo.
61. The method of claim 53, wherein the composition is selected from the group  
5 consisting of a gel, cream, paste, lotion, spray, suspension, dispersion, salve,  
hydrogel, ointment, and a biocompatible matrix.
62. A pharmaceutical composition comprising wound healing polypeptide  
comprising the amino acid sequence LKKTET and conservative variants thereof  
having wound healing activity, and a pharmaceutically acceptable carrier.
- 10 63 The pharmaceutical composition of claim 62, wherein the wound healing  
polypeptide is thymosin  $\beta$ 4 or an isoform of thymosin  $\beta$ 4.
64. The pharmaceutical composition of claim 62 in a controlled release formulation.
65. The pharmaceutical composition of claim 62 in a liposomal form.
66. The pharmaceutical composition of claim 62 in a lyophilized form.
- 15 67. The pharmaceutical composition of claim 62 in a unit dosage form.

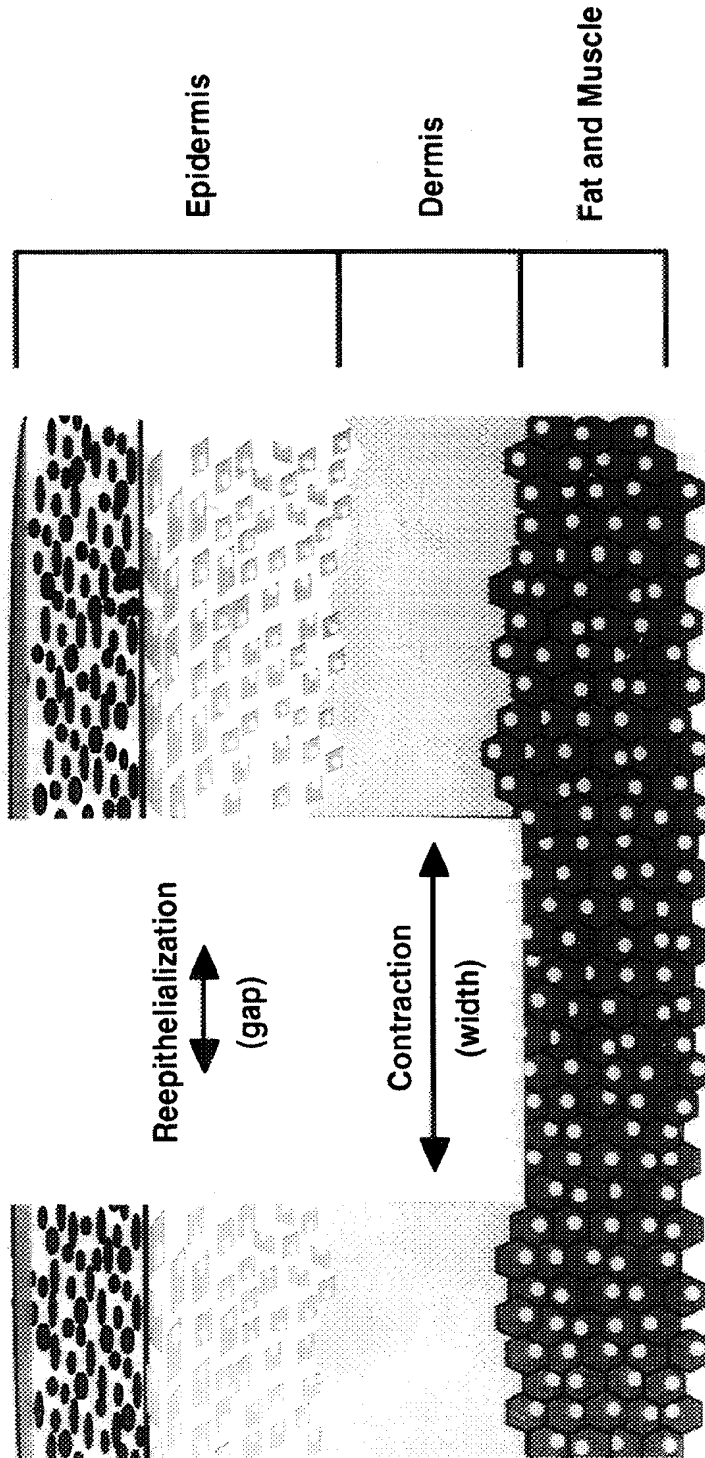


FIG. 1

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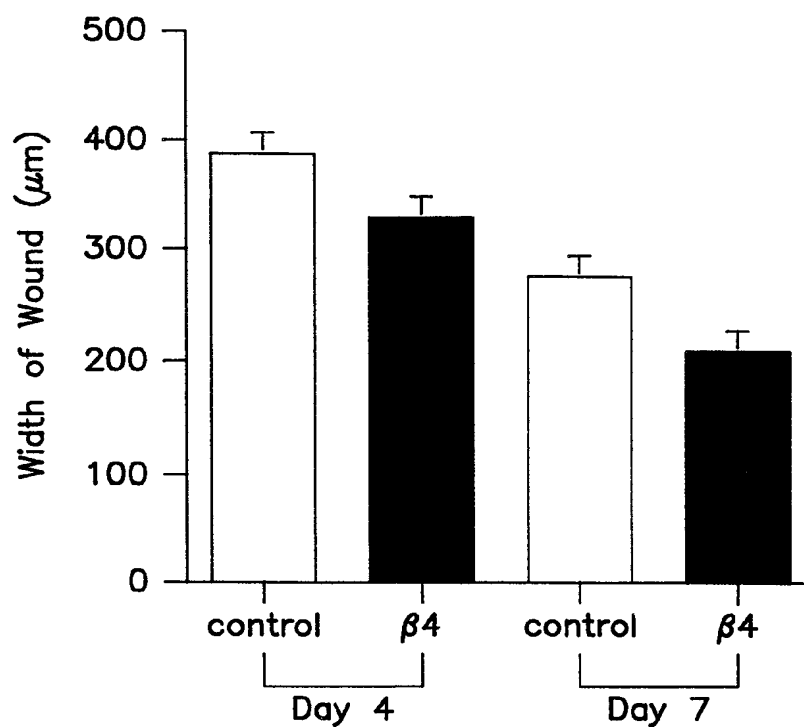


FIG. 2A

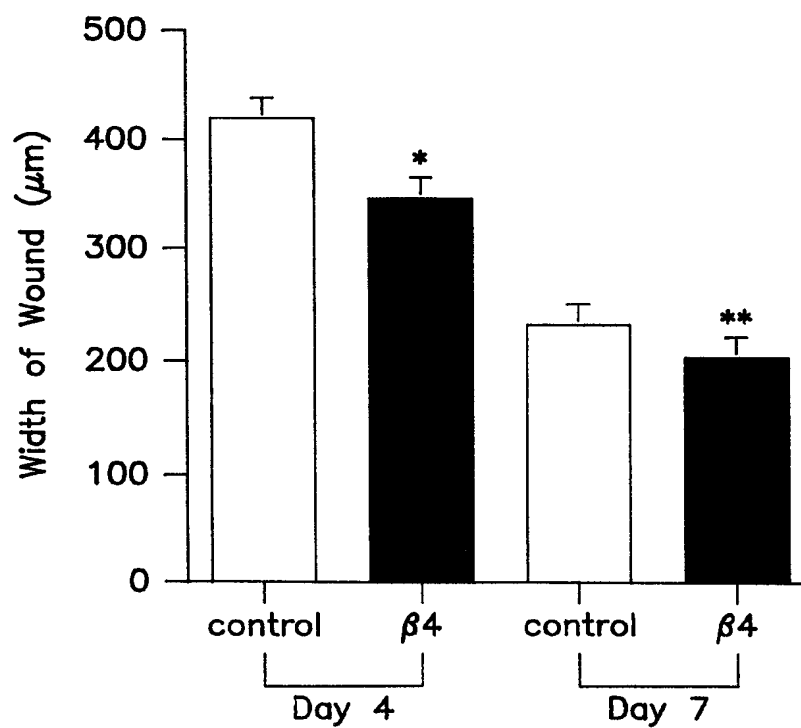


FIG. 2B

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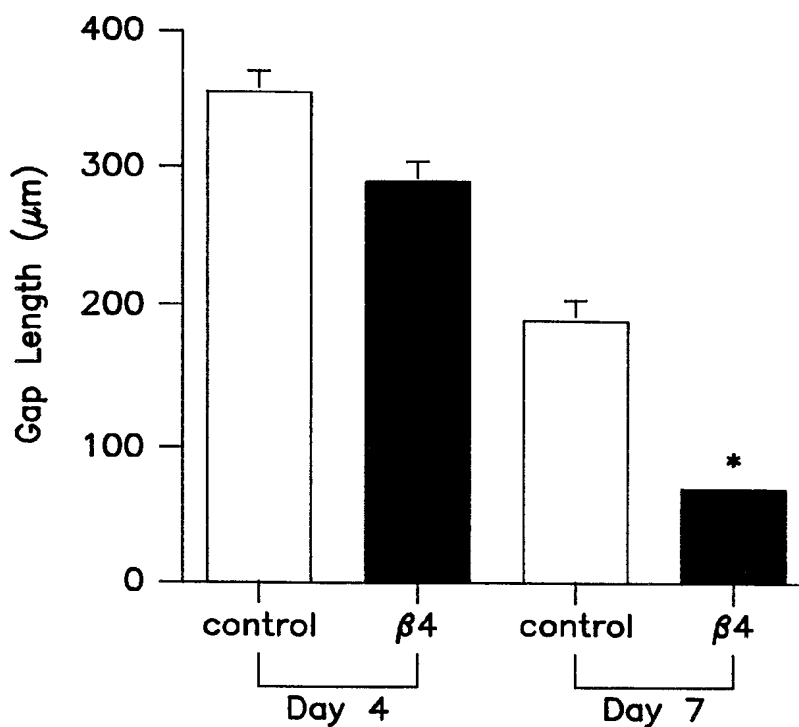


FIG. 3A

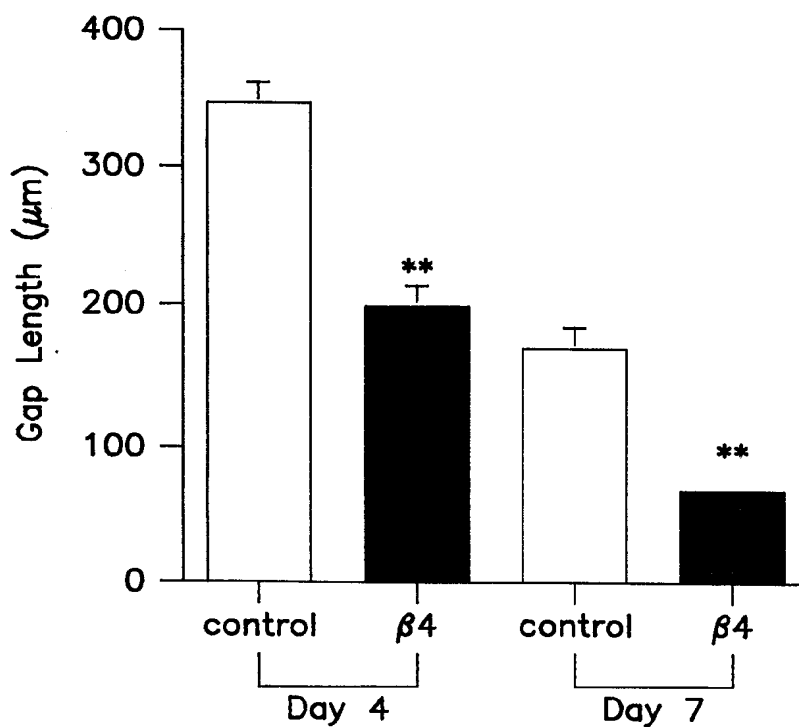


FIG. 3B

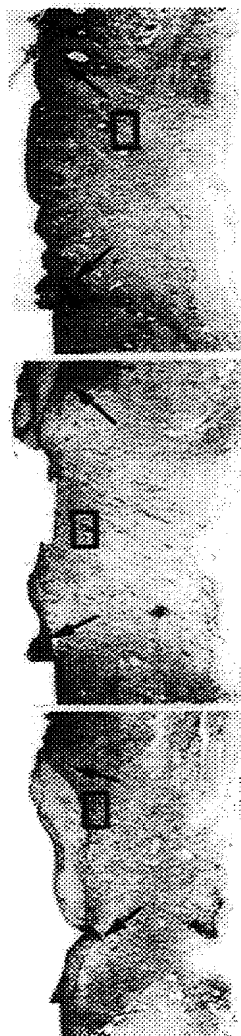


FIG. 4c

FIG. 4b

FIG. 4a



FIG. 4f

FIG. 4e

FIG. 4d



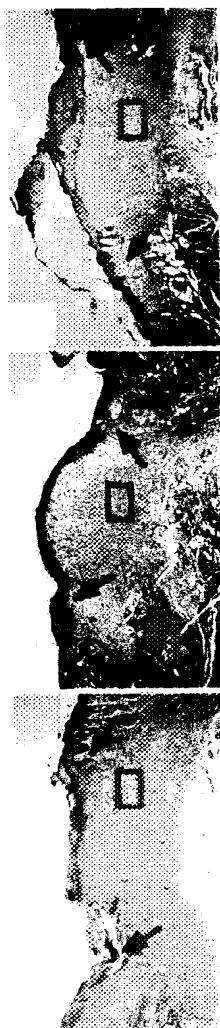


FIG. 5c

FIG. 5b

FIG. 5a



FIG. 5f

FIG. 5e

FIG. 5d

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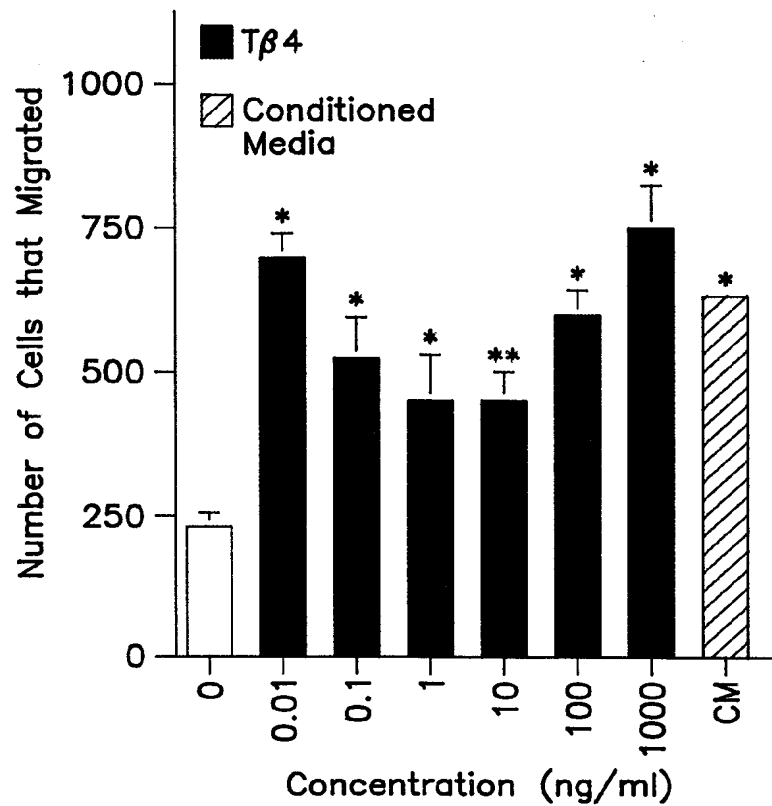


FIG. 6

Thymosin  $\beta$ 4 Stimulates  
Migration of Human Corneal  
Epithelial Cells

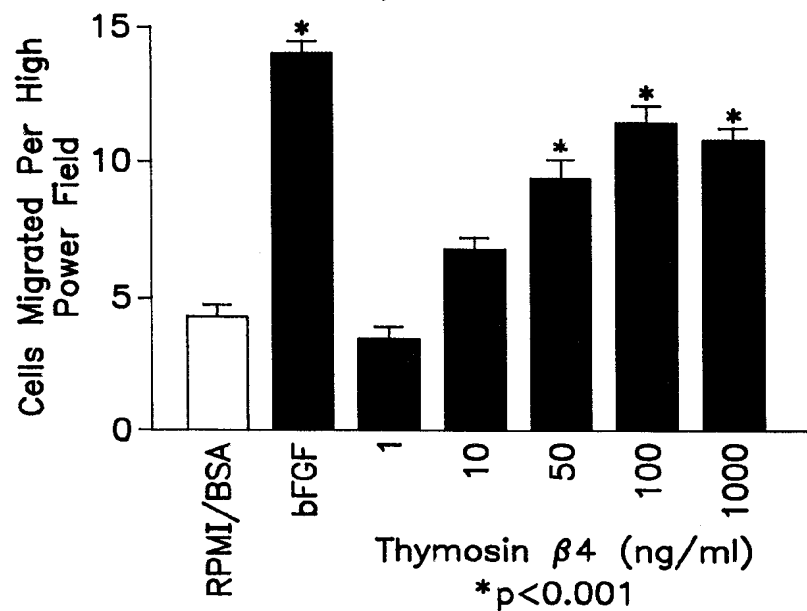
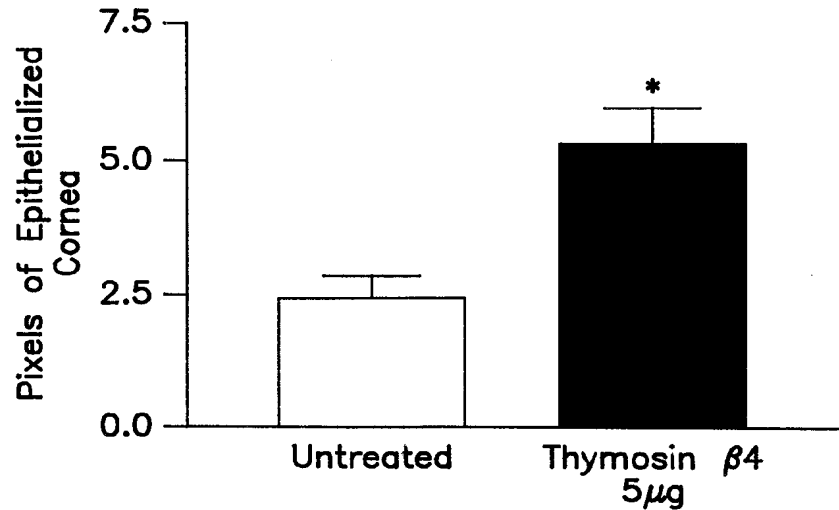


FIG. 7

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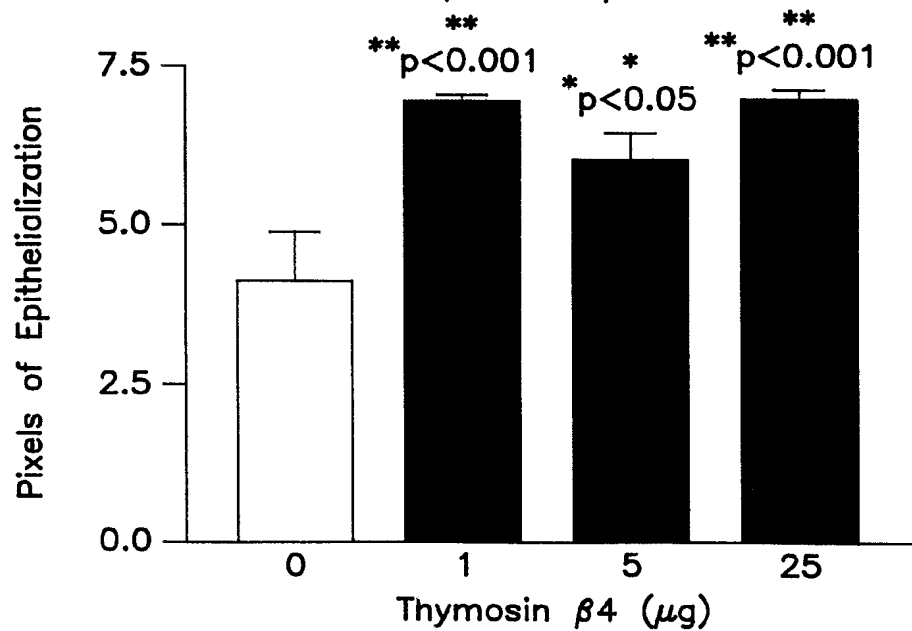
Thymosin  $\beta 4$  Stimulates  
Corneal Re-epithelialization in  
the Rat Cornea at 24 Hours

\*  $p=0.003$ 

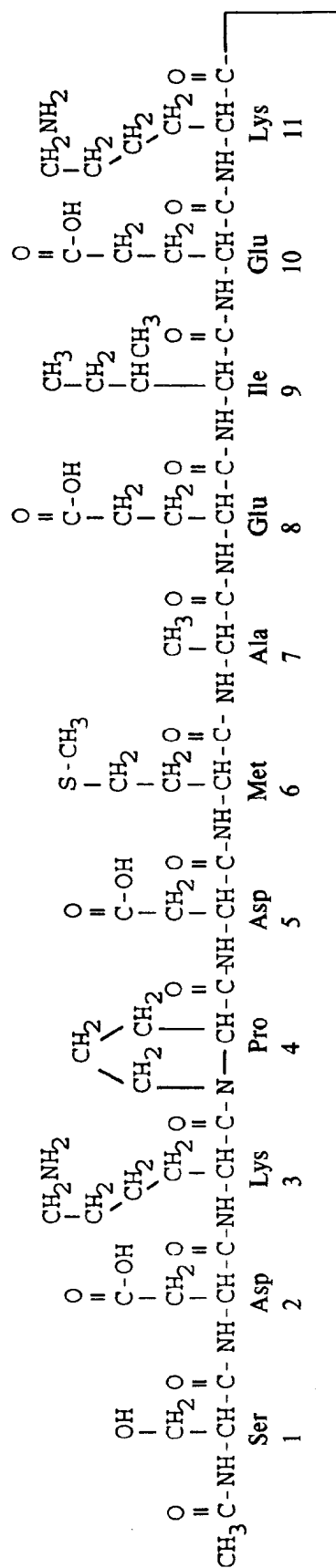
n=6

**FIG. 8**

Thymosin  $\beta 4$  Stimulates  
Re-epithelialization in the Rat  
Cornea at 24 Hours:  
Dose Response Experiment

**FIG. 9**

## Structural Formula of Thymosin Beta 4



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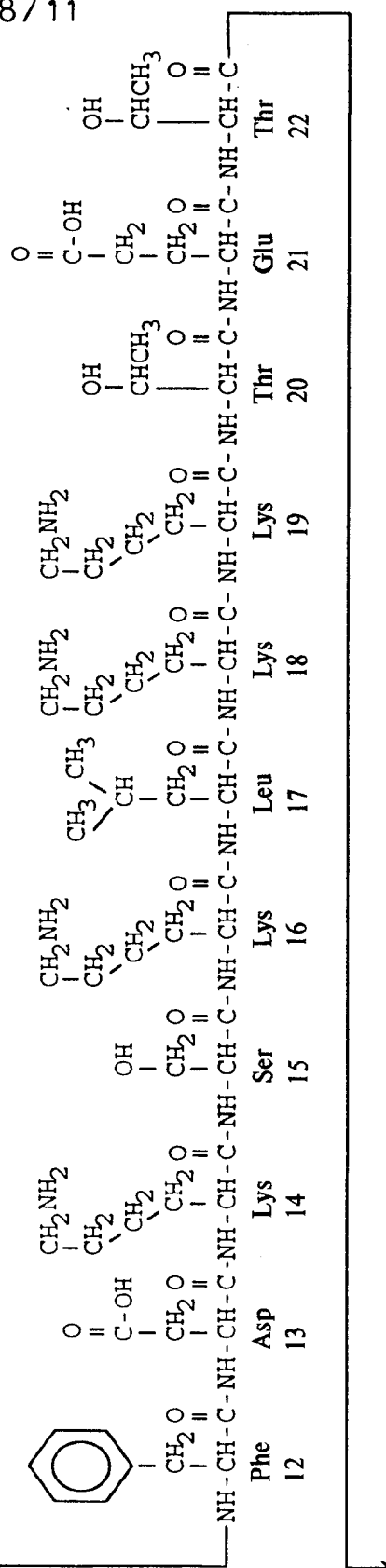


FIG. 10a

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Structural Formula of Thymosin Beta 4

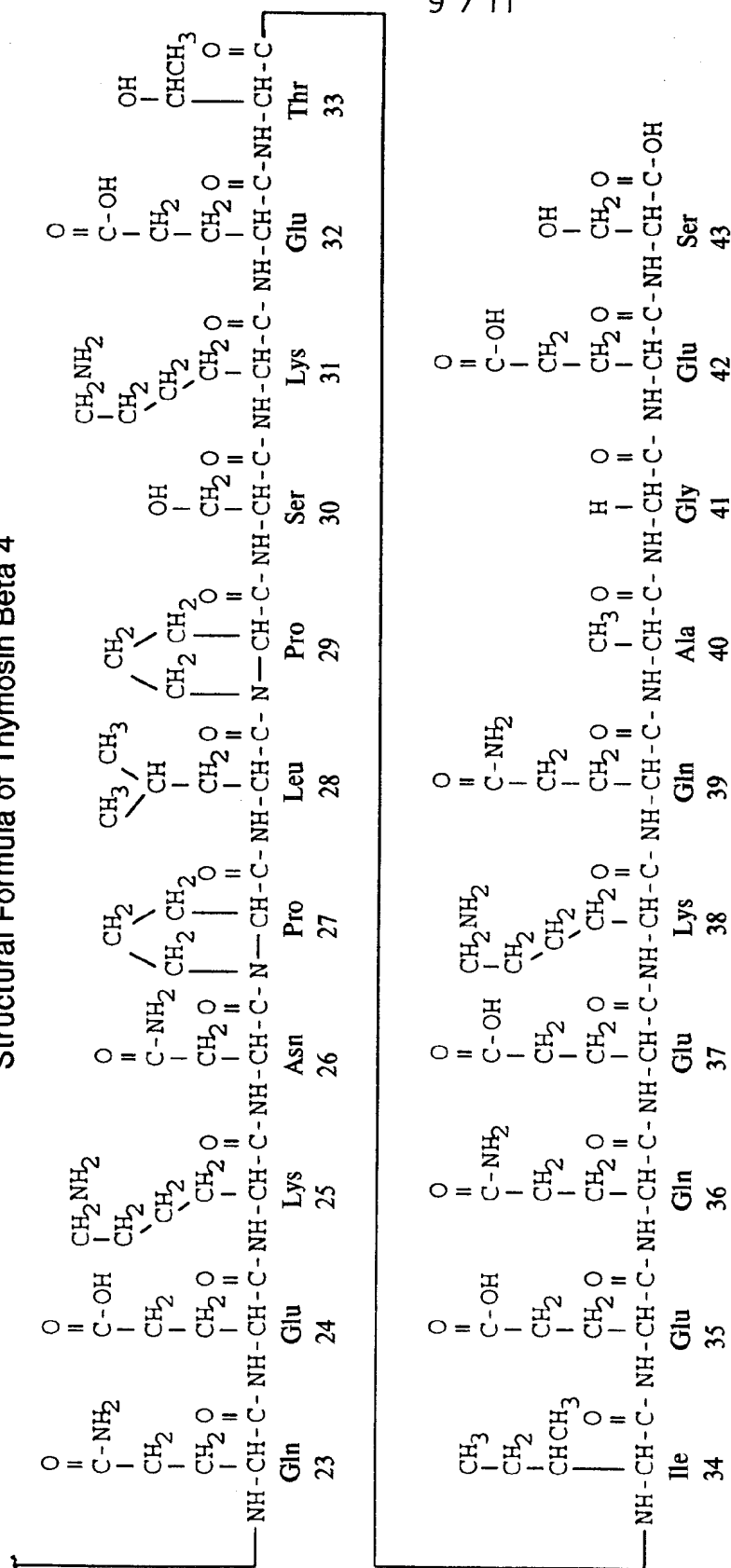


FIG. 10b

Amino Acid Sequence of Thymosin $\beta_4$ and other $\beta$ -Thymosins		5	10	15	20	25	30	35	40
		.....HELIX.....					....HELIX....		
$T\beta_4$	ac-SDKP DMAEI EKFDK SKLKK TETQE KNPLP SKETI EQEDQ AGES								
$T\beta_4^{Ala}$	ac-AKDP DMAEI EKFDK SKLKK TETQE KNPLP SKETI EQEKQ AGES								
$T\beta_4^{Xen}$	ac-SDKP DMAEI EKFDK AKLKK TETQE KNPLP SKETI EQEKQ STES								
$T\beta_9$	ac-ADKP DLGEI NSFDK AKLKK TETQE KNPLP TKETI EQEKQ AK								
$T\beta_9^{Met}$	ac-ADKP DMGEI NSFDK AKLKK TETQE KNPLP TKETI EQEKQ AK								
$T\beta_{10}$	ac-ADKP DMGEI ASFDK AKLKK TETQE KNPLP TKETI EQEKQ SEIS								
$T\beta_{11}$	ac-SDKP NLEEV ASFDK TKLKK TETQE KNPLP TKETI EQEKQ AS								
$T\beta_{12}$	ac-SDKP DLAEV SNFDK TKLKK TETQE KNPLP TKETI EQEKQ ATA								
$T\beta_{12}^{perch}$	ac-SDKP DISEV TSFDK TKLKK TETQE KNPLP SKETI EQEKA AATS								
$T\beta_{13}$	ac-ADKP DMGEI ASFDK AKLKK TETQE KNPLP TKETI EQEKQ AK								
$T\beta_{14}$	ac-SDKP DISEV SSFDK TKLKK TETAE KNPLP TKETI EQELT A								
$T\beta_{15}$	ac-SDKP DLSEV ETEFDK SKLKK TNTEE KNPLP SKETI QQEKE YNQRS								
$T\beta_{scallops}$	ac-SDKP FVSEV ANFDK SKLKK TETAE KNPLP TKETI QQEKE A								
$T\beta_{sea urch}$	ac-ADKP DVSEV STFDK SKLKK TETQE KNPLP TKDTI EQEKQ G								

FIG. 11a

### Phylogenetic Distribution of Thymosin $\beta_4$ -Like Peptides

Species	First peptide	Second peptide	Third peptide
Human	$\beta_4$	$\beta_{10}$	$\beta_{15}$
Rat, mouse, cat	$\beta_4$	$\beta_{10}$	$\beta_{15}$ (rat tumor)
Calf	$\beta_4$	$\beta_9$	
Pig, sheep	$\beta_4$	$\beta_9^{\text{Met}}$	
Horse, chicken, gecko	$\beta_4$		
<i>Xenopus laevis</i>	$\beta_4^{\text{Xen}}$		
Rainbow trout	$\beta_{11}$	$\beta_{12}$	
Perch	$\beta_{12}^{\text{perch}}$		
Whale	$\beta_{13}$		
Sea urchin	$\beta_{14}$	$\beta^{\text{sea urchin}}$	
Scallop	$\beta^{\text{scallop}}$		

FIG. 11b

# INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 99/17282

## A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 A61K38/22 A61K39/395

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 A61K C07K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	MALINDA K M ET AL: "Thymosin beta 4 stimulates directional migration of human umbilical vein endothelial cells." FASEB JOURNAL, vol. 11, no. 6, May 1997 (1997-05), pages 474-481, XP002125695	1,2,5-8, 13, 16-19, 23-28, 31-35, 38,53, 57-64
Y	the whole document	3,4,11, 12,14, 15,22, 29,30, 36,37
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☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

\* Special categories of cited documents :

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

Date of the actual completion of the international search

17 December 1999

Date of mailing of the international search report

12/01/2000

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Teyssier, B



# INTERNATIONAL SEARCH REPORT

Inter- national Application No  
PCT/US 99/17282

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	SUN H -Q ET AL: "BETA-THYMOSINS ARE NOT SIMPLE ACTIN MONOMER BUFFERING PROTEINS. INSIGHTS FROM OVEREXPRESSION STUDIES" JOURNAL OF BIOLOGICAL CHEMISTRY, vol. 271, no. 16, April 1996 (1996-04), pages 9223-9230, XP002041936 ISSN: 0021-9258 page 9223, column 1, line 15 - line 18 page 9229; figure 3	1,2, 8-10,13, 19-21, 23,24, 26,53, 60,62,63
X	page 9224, section "Quantitative Immunoblotting"	45,46
Y	page 9227, column 1, line 5 -page 9228, column 1, line 3 ---	11,22,36
Y	NIMNI M E: "Polypeptide growth factors: targeted delivery systems" BIOMATERIALS, vol. 18, no. 18, 1997, pages 1201-1225, XP004086390 ISSN: 0142-9612 page 1210, column 1, line 15 - line 53 page 1203 -page 1211	3,4,11, 12,14, 15,22, 29,30, 36,37
A	page 1205, section "Keloid and hypertrophic scars"	41-44
A	pages 1215-1216, sections "Delivery of growth factors in wound healing" and "Some novel and potentially useful approaches for local delivery of growth factors"	38-40, 54-61,64
Y	WO 96 16983 A (JOLLA CANCER RES FOUND) 6 June 1996 (1996-06-06) page 1 -page 2; claims 1,4	11,12, 22,36,37
A	claims 5,6,8,11,16-19 ---	6,7,17, 18,33, 34,58, 61,64
Y	FRANK, S ET AL: "Regulation of vascular endothelial growth factor expression in cultured keratinocytes and implications for normal and impaired wound healing" J. BIOL. CHEM., vol. 270, no. 21, May 1995 (1995-05), pages 12607-12613, XP002125696 page 12607, column 2, line 13 - line 26 page 12609, column 2, line 7 - line 12 ---	11,12, 22,36,37
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# INTERNATIONAL SEARCH REPORT

Inter. Application No  
PCT/US 99/17282

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 97 48805 A (CHILDRENS MEDICAL CENTER) 24 December 1997 (1997-12-24) page 14, line 17 -page 16, line 17	45,46
X	page 11, line 20 - line 22	53
A	page 16, line 19 -page 23, line 9	41-44, 47-49
X	EP 0 124 779 A (UNIV GEORGE WASHINGTON) 14 November 1984 (1984-11-14) claims	45,46
A	WO 94 11499 A (MAX PLANCK GESELLSCHAFT) 26 May 1994 (1994-05-26) claims 18-24	50-52
A	SABOLINSKI M L ET AL: "Cultured skin as a 'smart material' for healing wounds: experience in venous ulcers" BIOMATERIALS, vol. 17, no. 3, 1996, pages 311-320, XP004032797 ISSN: 0142-9612 Introduction	26
A	HANNAPPEL E & WARTENBERG F: "Actin-sequestering ability of thymosin beta 4, thymosin beta 4 fragments and thymosin beta 4-like peptides as assessed by the DNase I inhibition assay" BIOL. CHEM., vol. 374, February 1993 (1993-02), pages 117-122, XP002118663 page 121, column 1, line 8 -column 2, line 17	1,2,9, 10,20, 21,23, 24,62,63

## INTERNATIONAL SEARCH REPORT

International application No.

PCT/US 99/17282

### Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.:  
because they relate to subject matter not required to be searched by this Authority, namely:  
  
see FURTHER INFORMATION sheet PCT/ISA/210
2. ☐ Claims Nos.:  
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
3. ☐ Claims Nos.:  
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

### Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

#### Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

**FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210**

Continuation of Box I.1

Although claims 1-25, 27-44, 47-49, 53-59 and 61 are directed to a method of treatment of the human/animal body and claims 45 and 46 to a diagnostic method practised on the human/animal body, the search has been carried out and based on the alleged effects of the compound/composition.

# INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 99/17282

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9616983 A	06-06-1996	US 5654267 A AU 4412396 A CA 2206175 A EP 0797584 A JP 10509980 T US 5830504 A	05-08-1997 19-06-1996 06-06-1996 01-10-1997 29-09-1998 03-11-1998
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